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- ROSTRUM® 5% - Injectable Solution
- ROSTRUM® LARGE BREED - Oral Tablet
- ROSTRUM® PLUS - Otic Emulsion
- RUMITEN® Oral suspension
- SENILPET® CEREBRAL 20 Oral tablet.
- SENILPET® CEREBRAL 5 Oral tablet.
- SILIMADRAG® Oral suspension.
- SINPUL® Shampoo
- SINPUL® SPRAY Solution
- SINPULDRY® CAT - Topical powder
- SINPULDRY® DOG - Topical powder
- SINPULKILL® CATS - Collar

- SINPULKILL® Dog Collar
- SINPULSPOT® PLUS Large Breed - Spot on
- SINPULSPOT® PLUS Medium Breed - Spot on
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- SIR DOG® Black - Shampoo
- SIR DOG® CONDITIONER
- SIR DOG® ODOR CONTROL - Shampoo
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- SIR DOG® SHED CONTROL - Shampoo
- SIR DOG® WHITE - Shampoo
- SKINDRAG® Ceramides - Shampoo
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- SKINDRAG® Vitamin E - Conditioner
- SUAVIPET® Topic solution
- SUPERPET® OMEGA ADULT DOG - Oral Solution
- SUPERPET® OMEGA CAT - Oral Solution
- SUPERPET® OMEGA PUPPY - Oral Solution
- SUPERPET® OMEGA SENIOR - Oral Solution
- SUPOLEN® Oral tablet
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- TRANSIMED® Otic and Topical Suspension
- TRIAMCOL® Injectable solution
- TRIPLE- Injectable solution
- ULTRAFIL® PLUS Otic Suspension
- URSOVET® Oral suspension
- VERMIQUANTREL® Oral tablet
- VIDATOL® Injectable solution.
- XILA-10® - Injectable solution

ACTIGOR® INJECTABLE SOLUTION.

SOLUCIÓN INYECTABLE.

TÓNICO FOSFORADO + OLIGOELEMENTOS.

Technical Specification

SPECIES

Bovines, ovine, goats, pigs and horses.

Recommended to compensate diets low in phosphorus and trace- elements to balance the demands of pregnancy, growing up, reproduction, lactation, intense exercise, convalescence and improve the appearance (fur coat).

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Phosphorus tonic + Trace elements.

COMPOSITION

Each 100 mL of injectable solution contains:
Sodium Glycerol phosphate x 5.5 H2O.....28.49 g
(equivalent to 2.8 g of phosphorus)
Cobalt Chloride 6 H2O.....0.004 g
(equivalent to 1 mg de Cobalt)
Ammonium Molybdate x 4 H2O.....0.0092 g
(equivalent to 5 mg of Molybdenum)
Sodium selenite x 5 H2O.....0.0333 g
(equivalent to 10 mg of Selenium)
Zinc Sulfate x 7 H2O.....0.110 g
(equivalent to 25 mg of Zinc)
Manganese Sulfate x H2O.....0.077 g
(equivalent to 25 mg of Manganese)
Nicotinic acid.....0.500 g
Excipients q.s.p.....100 mL

ROUTE OD ADMINISTRATION AND DOSAGE

Administration: Intravenous, subcutaneous and intramuscular.

Dose:

It's suggested:

- Horses and bovines, adults: 10 - 20 mL, IV o IM.
- Horses and bovines, young: 3 - 5 mL, IV o IM.
- Pigs, adults: 3 - 6 mL, IM.
- Pigs, young: 1 - 1,5 mL, IM.
- Ovine and goats, adults: 2 - 4 mL, IM o SC.
- Ovine and goats, young: 0,5 - 1 mL, IM.

Administer as one single dose.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

None.

CONSERVATION

Keep at room temperature between 15 and 30ºC, protected from light.

CONDITION OF SALE

Sale with Veterinarian doctor prescription only.

PRESENTATION

20 mL and 100 mL vial

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 1270
- Dominican Republic: Reg. No. 5609
- El Salvador: VE2013094807

COUNTRIES WHERE IT IS MARKETED

Distribution in El Salvador:
Rafael Alfredo Alfaro Castillo.
8a C. Pte. Pje. Moreno N° 112, Col. Flor Blanca.
San Salvador - El Salvador.

AIR PROFOUNDLY® ORAL SOLUTION

SOLUCIÓN ORAL.

AYUDA A DESPEJAR LAS VÍAS RESPIRATORIAS FAVORECIENDO UNA MAYOR CAPACIDAD AERÓBICA.

Technical Specification

SPECIES

Horses.

Help clearing the airways by favoring a greater aerobic capacity, specially indicated for horses in training or subjected to strenuous physical activity. Indicated for high effort competition animals, like leaping, polo, endure, full test, rodeo, etc.

DOSAGE FORM

Oral solution.

THERAPEUTIC ACTION

Helps clear the airways favoring greater aerobic capacity.

COMPOSITION

Contain essential oils of *Menthae piperitae* and *Eucaliptus globulus* from: menthol, mentone, flavonoids, phenolic acid, tannins, triterpene lactone, cineole, eucalyptol, terpineol, alpha-pinene.

INGREDIENTS

Purified water, Glycerin, Propylene Glycol, Polyoxyl 40 Hydrogenated Castor Oil, 2-Pyrrolidone, Eucalyptus Oil, Peppermint Oil, Sodium Carboxymethylcellulose, Podium Bisulfite and authorized preservatives.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration.

- Moderate exercise: 15 mL
- Intense exercise: 30 mL

WARNINGS

- Mantener fuera del alcance de los niños.
- Agitar antes de usar.
- Mantener bien cerrado.

OBSERVATIONS

"EXCLUSIVE USE IN ANIMAL FEED"
"IT DOES NOT CORRESPOND TO A COMPLETE FOOD"

CONSERVATION

- Store in a cool dry place.
- Keep it tightly closed.

PRESENTATION

In a 1 liter bottle.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. LENAA N° RM 03-008N

ALERDRAG® SHAMPOO

SHAMPOO.

ANTIALÉRGICO DE USO TÓPICO.

Technical Specification

SPECIES

Dogs and cats.

Indicated for treatment of allergic dermatitis symptoms, such as pruritus and inflammation.

DOSAGE FORM

Shampoo.

THERAPEUTIC ACTION

Topical anti-allergic product.

COMPOSITION

Each 100 mL contains:
Hydrocortisone Acetate.....0.5581 g
(Equivalent to 0.5 g of Hydrocortisone base)
Excipients q.s.p.....100 mL

INDICATIONS

- Do not use in animals with purulent skin infections or open or very serious wounds.
- Do not administer to animals hypersensitive to Hydrocortisone, who have Cushing's Syndrome.
- Do not administer to animals younger than 7 months of age.
- Do not administer to pregnant or lactating females.

MODE OF APPLICATION

- Wet pet’s coat thoroughly with abundant water.
- Apply Alerdrag in a sufficient amount.
- Massage and let stand in for 10 to 15 minutes and rinse with plenty of water.
- Repeat the bath 3 times per week, or according the instructions given by your Veterinarian Doctor.

DRUG INTERACTIONS

In case of prolonged therapy or in large areas of skin, avoid concomitant administration with Amphotericin B or caliuretic diuretics (Furosemide, Thiazides) as hypokalemia may occur; Cyclosporine because the blood levels of both drugs can be increased, with mutual inhibition of liver metabolism; with ulcerogenic drugs (eg non-steroidal anti-inflammatory drugs), since the risk of gastrointestinal ulceration may be increased. Patients treated with corticosteroids at immunosuppressive doses generally should not receive live attenuated virus vaccines, because viral replication may be enhanced.

CONTRAINDICATIONS

- Do not use in animals with purulent skin infections or open or very serious wounds.
- Do not administer to animals hypersensitive to Hydrocortisone, who have Cushing's Syndrome.
- Do not administer to animals younger than 7 months of age.
- Do not administer to pregnant or lactating females.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Avoid inhalation, ingestion or direct contact with the use of gloves and mask. Wash hands thoroughly after use.
- Do not handle by people hypersensitive to Hydrocortisone.
- Do not handle by pregnant women.

WARNINGS

Advertencias y precauciones especiales de uso:

- No utilizar este producto sin previo diagnóstico.
- En caso de causar irritación, se debería descontinuar su uso.
- En caso de infecciones bacterianas, micóticas o parasitarias, se debe dar el tratamiento adecuado con indicación Médico Veterinaria.
- Uso externo.
- Mantener fuera del alcance de los niños.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:

- Empty containers can be discarded as household waste, without any special precautions.
- Do not dispose of containers with product residues on the ground or water courses.
- For expired or unused products contact the manufacturer laboratory.

CONSERVATION

Store between 15 and 30 ° C, protected from light.

CONDITION OF SALE

To be supplied only on veterinary prescription.
For Peru: Free Sale

PRESENTATION

A bottle containing 150 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 0147
- Costa Rica: Reg. No.MAG CL4-45-04-4310
- El Salvador: VE2015105120
- Panama: Reg. No. RF-4182-08
- Bolivia Reg. SENASAG PUV-F-N° 006143/14
- Rep. Dominicana: Reg. Nº 8669

APETICAT® SYRUP

JARABE.

SUPLEMENTO VITAMÍNICO CON TAURINA - CARNITINA PARA GATOS.

Technical Specification

SPECIES

Cats.

Apeticat® is a supplement specially formulated for cats based on Taurine and Carnitine, supplemented with amino acids, vitamins and Calcium. Its formula increases your pet's desire to eat, acting as a bioenergizer and an enhancer of the body's metabolic functions. Taurine is an amino acid that cats only synthesize in very small amounts, not enough to fully satisfy the physiological needs of the animal, its presence in adequate quantity is essential in the diet to maintain the health of your pet.

DOSAGE FORM

Syrup.

THERAPEUTIC ACTION

Vitamin supplement with Taurine - Carnitine for cats.

COMPOSITION

Each 100 mL contains:
Carnitine HCl.....1.6 g
Sorbitol.....25 g
Taurine.....1 g
Choline Chloride.....9 mg
Vitamin B1.....5 mg
Vitamin B2.....4 mg
Vitamin B6.....4 mg
Vitamin B12.....20 µg
Nicotinic acid.....6 mg
Calcium Pantothenate.....5 mg
Special excipients q.s.p.....100 mL

INGREDIENTS

Declared vitamins and minerals, Taurine, purified water, meat essence, salmon essence and authorized preservatives.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer orally 1 teaspoon (5 mL) for every 4 Kg of weight.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

DOES NOT CONSTITUTE A COMPLETE FOOD
VETERINARY USE
EXCLUSIVE USE IN ANIMAL FEED

CONSERVATION

Store in a cool, dry place, protected from light, below 30ºC.

CONDITION OF SALE

Direct sale.

PRESENTATION

A bottle containing 100 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. LENAA N°: RM 03-008N

APETIPET® SYRUP

JARABE.

SUPLEMENTO VITAMÍNICO PARA PERROS.

Technical Specification

SPECIES

Dogs.

Apetipet is a supplement based on Carnitine, complemented with amino acids, vitamins and calcium. Its formula increases your pet's appetite.

DOSAGE FORM

Syrup.

THERAPEUTIC ACTION

Vitamin supplement for dogs.

COMPOSITION

Each 100 mL contains:
Carnitine HCl.....5 g
Sorbitol.....25 g
Methionine.....1 g
Choline Chloride.....1 g
Vitamin B1.....15 mg
Vitamin B2.....5 mg
Vitamin B6.....15 mg
Vitamin B12.....0.28 mg
Nicotinic acid.....25 mg
Calcium Pantothenate.....10 mg
Excipients q.s.p.....100 mL

INGREDIENTS

Purified water, Sorbitol, Carnitine HCl, Methionine, Declared Vitamins and Minerals, Meat Essence and authorized preservatives.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer orally, 1 teaspoon (5 mL) for every 10 Kg of weight.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

DOES NOT CONSTITUTE A COMPLETE FOOD
VETERINARY USE
EXCLUSIVE USE IN ANIMAL FEED

CONSERVATION

Store in a cool, dry place, protected from light, below 30ºC

CONDITION OF SALE

Direct sale.

PRESENTATION

A bottle containing 100 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG RM 03-008N

ARTRIOFIN® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANTIINFLAMATORIO NO ESTEROIDAL

Technical Specification

SPECIES

Dogs

Indicated in the relief of pain and inflammation in dogs. It is especially indicated in the relief of signs associated with canine osteoarthritis. It is also recommended in the control of post-operative pain associated with orthopedic and soft tissue surgeries (including intraocular surgery).

DOSAGE FORM

Injectable solution

THERAPEUTIC ACTION

Non-steroidal anti-inflammatory

COMPOSITION

Each 1 mL of injectable solution contains:
Carprofen.....44 mg
Excipients q.s.p.....1 mL

INDICATIONS

- Do not administer in animals with hypersensitivity to Carprofen.
- Do not administer intramuscularly.
- Do not exceed the indicated dose.

EFFECTIVENESS

Period of effectiveness: 36 months.

ROUTE OD ADMINISTRATION AND DOSAGE

Subcutaneous administration

- **Active ingredient dose**: 4.4 mg / kg once a day or 2.2 mg / kg twice a day, for up to 5 days.
- **Product dose:** 1 mL for every 10 kilos of weight once a day, or divided into two 0.5 mL doses for every 10 kilos of weight, every 12 hours, for up to 5 days.

In the control of post-operative pain, its administration is recommended 2 hours before the procedure.

CONTRAINDICATIONS

- Do not administer in animals with hypersensitivity to Carprofen.
- Do not administer intramuscularly.
- Do not exceed the indicated dose.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

Long-term use of **Artriofin®** can cause gastrointestinal injury, loss of appetite, vomiting and diarrhea.

CONSERVATION

Store in a cool, dry place, at room temperature between 15 and 30ºC.
Once the container is opened, use within 12 weeks. Discard the unused product after that period.

CONDITION OF SALE

Sale under Veterinary Medical prescription.

PRESENTATION

50 mL ampoule bottle

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG Nº 1801
Uruguay: Reg. Nº MGAP A-4456
Perú: Registro SENASA F.99.01.I.0076

AVAILABLE FOR SALE

Imported and Distributed in Uruguay by:
VIVAFIL S.A. RIO NEGRO 1107 Montevideo
Uruguay,
TEL 29001112
grupotecnovet@gmail.com
Director Técnico: DMTV Diego Cuadrado.

Imported and Distributed in Peru by:
Representaciones Durand SAC.
Av. Manuel Olguín N° 501 Oficina N° 604
Santiago de Surco Lima.

ARTRIOFIN® ORAL TABLETS.

COMPRIMIDO ORAL.

ANTIINFLAMATORIO NO ESTEROIDAL.

Technical Specification

SPECIES

Dogs.

Indicated in the relief of pain and inflammation associated with acute or chronic osteoarticular conditions, such as canine osteoarthritis. Its use is also recommended in the control of post-operative pain associated with orthopedic and soft tissue surgeries.

DOSAGE FORM

Oral tablets.

THERAPEUTIC ACTION

Nonsteroidal anti-inflammatory

COMPOSITION

Each tablet contains:
Carprofen..... 88 mg
Excipients q.s.p.....1 tablet

INDICATIONS

- Do not use with any other nonsteroidal anti-inflammatory product.
- Do not administer to pregnant or nursing females.
- Do not administer to dehydrated, hypovolemic or with hypotension, with heart, liver or kidney condition dogs.
- Do not administer to dogs with gastroduodenal ulcers, blood dyscrasia and hypersensitivity to the active ingredient or to other derivatives of propionic acid.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration.

- **Active ingredient dose:** 4.4 mg/Kg once a day or 2.2 mg/Kg twice a day, for 7 to 10 days.
- **Product dose:** 1 tablet for each 20 kg of body weight, once a day or divided in two doses of half tablet every 12 hours, for 7 to 10 days.

  The Veterinary Doctor must evaluate the patient to continue with the treatment. In the control of post-operative pain, its administration is recommended 2 hours before the procedure.

CONTRAINDICATIONS

- Do not use with any other nonsteroidal anti-inflammatory product.
- Do not administer to pregnant or nursing females.
- Do not administer to dehydrated, hypovolemic or with hypotension, with heart, liver or kidney condition dogs.
- Do not administer to dogs with gastroduodenal ulcers, blood dyscrasia and hypersensitivity to the active ingredient or to other derivatives of propionic acid.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

The long-term use of **Artriofin®** can cause gastrointestinal lesions, lack of appetite, vomiting and diarrhea. If any of these symptoms appears, ask immediately the treating Veterinarian Doctor.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° and 30°C.

CONDITION OF SALE

To be supplied only on veterinary prescription.

PRESENTATION

A box containing 10 tablets.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 1617
Costa Rica: Reg. N° MAG CL4-14-4-5800
Rep. Dominicana: Reg. Nº 8863
Uruguay: Reg. MGAP N° A-4457
Bolivia: Reg. SENASAG PUV- Nº 010001/21

AZANILVET® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

SEDANTE.

Technical Specification

SPECIES

Pigs.

Azanilvet® is a neuroleptic sedative especially indicated for pigs. Its use is recommended for regrouping, postpartum stress management, transport and anesthetic premedication of pigs.

- Regrouping: In fattening pigs and weaned piglets, the use of Azanilvet® is recommended 15 to 20 minutes prior to the regrouping of animals, since it prevents, delays and reduces the time and frequency of attacks up to 2 hours after application. In boars, the use of Azanilvet® 15 to 20 minutes prior to regrouping, allows reducing the intensity and violence of the aggressions for at least 4 hours after administration.
- Management of postpartum stress: The administration of Azanilvet® in sows at the time of delivery of the placenta, allows to control postpartum stress in the mother, thus improving the distribution of colostrum, reducing the incidence of diarrhea and obtaining a weaning weight in piglets from treated dams.
- Pig transport: The administration of Azanilvet® reduces mortality and weight loss in fattening pigs caused by transport stress.
- Premedication for anesthesia: The administration of Azanilvet® prior to anesthesia, allows obtaining a good sedation, achieving a better induction and anesthetic duration. Its effect is comparable or superior to that of other pre-anesthetics, allowing proper handling of the animal to perform various surgical procedures.

The sedative effect in pigs begins approximately 15 to 20 minutes after intramuscular administration, the effect remaining for 2 to 4 hours, depending on the dose used.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Sedative.

COMPOSITION

Each 1 mL of solution for injection contains:
Azaperone .....40 mg
Excipients q.s.p .....1 mL

INDICATIONS

- Do not administer intravenously, due to the presentation of a significant excitatory phase with cardiovascular effects.
- Do not administer in overexcited animals, as it increases the possibility of manifesting unwanted effects.
- Do not administer in pregnant females.

ROUTE OD ADMINISTRATION AND DOSAGE

Route of administration and method of use:
Administer deep intramuscularly using a long hypodermic needle, as close as possible to the back of the ear, perpendicular to the skin.

Dose:
Administer in a single dose. The dose varies according to the indication for use:
- Regrouping:
  Pigs and piglets: 1 mL / 20 Kg (equivalent to 2 mg / Kg).
  Boars: 0.5 mL / 20 Kg (equivalent to 1.0 mg / Kg).
- Postpartum stress management: 1 mL / 20 Kg (equivalent to 2 mg / Kg).
- Transportation of pigs: 0.2 to 0.3 mL / 20 Kg (equivalent to 0.4 to 0.6 mg / Kg).
- Premedication for anesthesia: 1 mL / 20 Kg (equivalent to 2 mg / Kg).

DRUG INTERACTIONS

Azaperone potentiates the action of general anesthetics, and the dose of these must be decreased when Azaperone is used as anesthetic premedication.

CONTRAINDICATIONS

- Do not administer intravenously, due to the presentation of a significant excitatory phase with cardiovascular effects.
- Do not administer in overexcited animals, as it increases the possibility of manifesting unwanted effects.
- Do not administer in pregnant females.

PRECAUTIONS

- Avoid use in very cold places, due to the risk of cardiovascular collapse secondary to peripheral vasodilation.
- Azaperone is not a substitute for proper anesthesia or analgesia.
- Azanilvet® should be administered in quiet environments to mitigate or eliminate possible unwanted effects.

SPECIAL PRECAUTIONS FOR USE

- Avoid use in very cold places, due to the risk of cardiovascular collapse secondary to peripheral vasodilation.
- Azaperone is not a substitute for proper anesthesia or analgesia.
- Azanilvet® should be administered in quiet environments to mitigate or eliminate possible unwanted effects.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Wash your hands after administering the product.
- In case of accidental self-injection, get immediate medical help.
- In case of contact with the skin and eyes, wash with plenty of water. If irritation develops and persists, see doctor.
- In case of ingestion, do not induce vomiting. Get medical help.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

In therapeutic doses, the use of Azaperone can produce hypothermia, hypotension and cardiorespiratory depression in pigs. Additionally, and usually in the presence of disturbing environments, other effects may occur such as: salivation, panting, stacking, muscle tremors, disorientation and excitement, which are transitory in nature.
In boars, therapeutic doses of Azaperone (1.0 mg / Kg) can cause temporary relaxation of the penis with eventual damage to it. This effect is more frequent in overdose.

GUARD PERIOD

Meat: 6 days.

OBSERVATIONS

Overdose:
Although unwanted effects may occur at therapeutic doses, Azaperone overdose added to administration in disturbing environments increases the probability of these effects occurring in a very marked way.
In boars, overdose can more frequently cause temporary relaxation of the penis, with eventual damage to the penis.

Special precautions for disposal of unused product or waste material:
Empty containers (without product) discard with household waste. In the case of expired product or containers with rest of the product, you should contact the manufacturing laboratory to receive recommendations for its correct disposal.

CONSERVATION

Storage conditions:
Keep the product at a temperature between 2 and 30 ºC, protected from light.
Once the container is opened, use within 4 weeks and must be stored at a temperature between 15 and 30ºC, protected from light. Discard the unused product after that period of time.

CONDITION OF SALE

Sale under Veterinary Medical prescription.

PRESENTATION

100 mL vial

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG Nº 2362
Costa Rica: Reg. MAG N° CL4-3-11-6534

BILIFAR® ORAL POWDER

POLVO ORAL.

ESTIMULANTE DE LA DIGESTIÓN RUMINAL

Technical Specification

SPECIES

Cattle, sheep and goats.

BILIFAR® Oral Powder, is a stimulant of ruminal digestion. Its use is recommended in pathologies in which it is necessary to stimulate ruminal fermentation and / or increase blood glucose levels, such as simple ruminal indigestion, alkalosis or ruminal rot, ketosis adjuvant and / or sudden changes in diet. BILIFAR® is also an alternative source of minerals for ruminants.

DOSAGE FORM

Oral powder

THERAPEUTIC ACTION

Ruminal Digestion Stimulant

COMPOSITION

Each bag with 120 grams contains:
Sodium propionate ..... 60 g
Calcium propionate ..... 40 g
Sodium Chloride ..... 18.5 g
Cobalt Chloride ..... 40 mg
Copper Sulfate ..... 150 mg
Manganese Sulfate ..... 200 mg
Iron Sulfate ..... 300 mg
Zinc Chloride ..... 10 mg
Excipients q.s.p ..... 120 grams

INDICATIONS

- Do not use in case of hypersensitivity to some components of the formula.
- Do not administer to dehydrated animals without prior hydration therapy.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration route:
Oral.

Product dosage:
- Adult cattle (cows, oxen, bulls) 450 to 600 Kg: 1 to 2 bags of 120 g dissolved in 1 to 3 liters of water.
- Young cattle (calves, steers) 150 to 300 Kg: 1/2 bag of 120 g dissolved in 1/2 to 1 liter of Water.
- Sheep and goats: 1/4 bag of 120 g dissolved in 1/4 liter of water.

The treatment can be repeated every 12 to 24 hours, for up to 10 days.

CONTRAINDICATIONS

- Do not use in case of hypersensitivity to some components of the formula.
- Do not administer to dehydrated animals without prior hydration therapy.

WARNINGS

- Mantener fuera del alcance de los niños.
- Respetar la dosis señalada considerando el margen de seguridad del Sulfato de Cobre.

GUARD PERIOD

Meat: 0 days.
Milk: 0 days.

OBSERVATIONS

Overdose:
Symptoms of Copper toxicity include hemolysis, hemoglobinuria, and jaundice. The oral toxic dose of Copper for sheep and calves is 20 to 110 mg / Kg; for cattle (adults) it is from 220 to 880 mg / Kg; and for goats it is 60 mg / Kg.

Special precautions for the disposal of unused product or waste material:
Eliminate the remains of unused product in its original container. Do not throw the empty container or with product remains, in rivers, lakes or torrents of water. Dispose of this product with caution with household waste.

CONSERVATION

Store in a cool, dry place, at room temperature between 15º and 30ºC. Use immediately once prepared and discard the excess product.

CONDITION OF SALE

Sale only with a Veterinary Medical prescription.

PRESENTATION

Sachet with 120 grams.

RECORDS

- Chile: Reg. SA. N° 280

BIO-POWER® EQUINE - ORAL PASTE

PASTA ORAL.

PROBIÓTICOS EN PASTA.

Technical Specification

SPECIES

Horses

Biopower® Equine is indicated in foals from birth, at weaning and under other stress conditions. Use in adult horses during transport, in periods of stress associated with training and competitions. It is also indicated as an aid in the treatment of enteric diseases and during recovery from surgery or injuries.
It is recommended to administer Biopower® Equine, concomitantly, in horses that are subjected to prolonged antimicrobial therapy, as an aid in the prevention of diarrhea due to bacterial overgrowth. Maintain the dosage of the product for a week after the antimicrobial therapy has been completed.

DOSAGE FORM

Oral Paste

THERAPEUTIC ACTION

Probiotics in paste

COMPOSITION

Each 30 g of pasta contains no less than:
Lactobacillus acidophilus ..... 2500 million u.f.c.
Bifidobacterium bifidum ..... 2500 million u.f.c.
Bacillus subtilis ..... 2500 million u.f.c.
Lactobacillus lactis ..... 2500 million u.f.c.

PROPERTIES

Due to its high concentration of beneficial microorganisms, Biopower Equino helps to correct the balance of the intestinal flora, favoring the inhibition of the growth of pathogenic organisms and providing digestive enzymes required in the digestion of food.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer orally, once a day for a week.

- **Adult horses:** 1 syringe x 30 grams.
- **Foals or young animals:** ½ syringe (15 grams).

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

- EXCLUSIVE USE IN ANIMAL FEED
- DOES NOT CORRESPOND TO A COMPLETE FOOD

CONSERVATION

Store in a cool, dry place, protected from light, at no more than 30ºC.

CONDITION OF SALE

OTC non prescription

PRESENTATION

Dosing syringe with 30 g

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. LENAA N°: RM 03-008N
Uruguay: Reg. MGAP N° 18850
Costa Rica: Lic. DAA-MAG 579-020

COUNTRIES WHERE IT IS MARKETED

Uruguay Importer:
VIVAFIL S.A.
RIO NEGRO 1107 Montevideo - Uruguay, TEL 29001112
grupotecnovet@gmail.com
Technical Director: DMTV Diego Cuadrado.

Imported and distributed in Costa Rica by:
Proventas de Cartago S.R.L.
100 meters east Hogares Crea, San Blas. Carthage. Tel: 2591 4624 Fax: 2591 5339

BIO-POWER® ORAL POWDER

POLVO ORAL.

PROBIÓTICOS.

Technical Specification

SPECIES

Dogs and cats.

- BIOPOWER® preserves and/or helps to recover an appropriate balance of normal gut flora in your pet.
- BIOPOWER® allows a better absorption of nutrients provided by food.
- BIOPOWER® provide microorganisms typical of digestive system, which are essential adjuvants to resolve gastrointestinal disorders of different origin, e.g. food changes and antibiotic therapy.
- In puppies, BIOPOWER® provides elements that favor the progressive establishment of normal intestinal flora.
- In senior animals, BIOPOWER® encourages the digestive function, preserving an appropriate stability of the intestinal flora of your pet.

DOSAGE FORM

Oral powder

THERAPEUTIC ACTION

Probiotics.

COMPOSITION

Each gram of powder contains:
Lactobacillus acidophilus..... 50,000,000 c.f.u.
Bifidobacterium bifidum.....50,000,000 c.f.u.
Enterococcus faecium.....50,000,000 c.f.u.
Excipients q.s.....1 g

INGREDIENTS

Lactobacillus acidophillus, Bifidobacterium bifidum, Enterococcus faecium, Sugar.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route. Administer once a day, directly into the animal's oral cavity.

- **Small dogs**: Dissolve 2.5 g (1/2 measuring spoon) in 10 mL of water (2 teaspoons)
- **Big dogs:** Dissolve 5.0 g (1 measuring spoon) in 15 mL of water (1 tablespoon)
- **Cats:** Dissolve 2.5 g (1/2 measuring spoon) in 10 mL of water (2 teaspoons)

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

- Includes measuring spoon.
- DOES NOT CONSTITUTE A COMPLETE FOOD
- EXCLUSIVE USE IN ANIMAL FEED

CONSERVATION

Store in a cool and dry place, out of direct light, below 30°C

PRESENTATION

100 g

PREPARED BY

Drag Pharma Chile Laboratory

RECORDS

Chile: Reg. LENAA N°: RM 03-008N
Bolivia: Reg. SENASAG PUV-A nº 008930/19

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

BOVIFORT® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANTIPARASITARIO DE AMPLIO ESPECTRO - FASCIOLICIDA.

Technical Specification

SPECIES

Bovines

Bovifort® is a broad-spectrum injectable antiparasitic that is indicated for the treatment of infestations by gastrointestinal, pulmonary and ectoparasites in cattle. Due to its specific action as a fasciolicide, it is also indicated in the treatment of hepatic dystomatosis, caused by adult Fasciola hepatica, also exercising adequate control over juvenile stages of 8 weeks or more. Its use is indicated in the treatment and control of the main internal and external parasites of bovines such as: Ostertagia sp .; Haemonchus sp .; Trichostrongylus sp .; Cooperia sp .; Oesophagostomum radiatum; Nematodirus sp .; Dictyocaulus viviparus; Fasciola hepatica; Haematopinus eurysternus; Linognathus vituli ,; Psoroptes bovis, Sarcoptes scabiei var bovis.

DOSAGE FORM

Injectable solution

THERAPEUTIC ACTION

Broad spectrum antiparasitic - Fasciolicida

COMPOSITION

Each 1 mL of solution for injection contains:
Ivermectin ..... 10 mg
Clorsulon ..... 100 mg
Excipients q.s.p ..... 1 mL

ROUTE OD ADMINISTRATION AND DOSAGE

Route of administration:
Subcutaneous.
Do not apply the product intravenously or intramuscularly.
In some cases there may be mild pain at the injection site.

Dosage of active ingredients:
Ivermectin: 0.2 mg / Kg and Chlorsulon: 2 mg / Kg
Single dose.

Product dosage:
1 mL for every 50 Kg of weight in a single dose.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Beef cattle: 49 days.

Milk cattle: Do not administer to animals whose milk is intended for human consumption.

CONSERVATION

Keep at room temperature between 15 ° and 30 ° C, in a dry place protected from light.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

50 mL ampoule bottle and 500 mL serum bottle

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 1880-B

CABATINA® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANTIHEMORRÁGICO.

Technical Specification

SPECIES

Dogs.

CABATINA® INJECTABLE is indicated for the initial emergency treatment of acute and over-acute poisonings generated by anticoagulant rodenticides.

DOSAGE FORM

Injectable solution

THERAPEUTIC ACTION

antihemorrhagic

COMPOSITION

Each 1 mL of solution for injection contains:
Phytomenadione.....10 mg
Excipients q.s.p..... 1 mL

PROPERTIES

Vitamin K is a cofactor in the last stage of hepatic synthesis of coagulation factors II (prothrombin), VII (proconvertin), IX (plasma thromboplastin component) and X (Stuart factor).

In veterinary medicine, the most frequent cause of the cessation of K-dependent coagulation factor formation is intoxication by anticoagulant rodenticides.

Phytomenadione (Vitamin K1) antagonizes the effects of anticoagulant rodenticidal agents and makes it possible to reactivate the production process of K-dependent coagulation factors.

INDICATIONS

- Do not administer in animals with known hypersensitivity to Phytomenadione (Vitamin K1).
- Do not administer in patients with severe liver failure.
- Do not administer intravenously due to the risk of anaphylactic reactions.
- Do not administer intramuscularly due to the risk of bleeding and bruising.
- Do not administer the product for more than 4 administrations due to the risk of producing hemolysis.

ROUTE OD ADMINISTRATION AND DOSAGE

Route of administration: Subcutaneous.

Dose:

- **Initial emergency treatment:**

Administer 2.5 to 5 mL of CABATINA® INJECTABLE product for every 10 kg of body weight (equivalent to 2.5 - 5 mg / kg of body weight of Phytomenadione, respectively), in a single dose.

According to the evolution of the clinical picture, type of rodenticide responsible for the intoxication and at the discretion of the treating Veterinarian, continue the treatment with another veterinary drug based on Phytomenadione administered orally.

- **Cases that require continuing with injectable treatment:**

In those cases that after the emergency dose, it is not possible to establish an oral therapy, treatment with the CABATINA® INJECTABLE product can be continued, in doses of 2.5 mL of product per 10 Kg of weight. body weight (equivalent to 2.5 mg / Kg of body weight of Phytomenadione), every 12 hours, with a maximum of 4 total administrations.

The duration of the injectable treatment (between 1 to 4 administrations) will depend on the type of rodenticide responsible for the poisoning, the evolution of the clinical picture and the criteria of the treating Veterinarian. Subsequently, treatment with another veterinary drug based on Phytomenadione can be continued orally.

How to use:
Administer the dose of the product at several injection sites to accelerate its absorption and using the smallest gauge possible to minimize the risk of bleeding.

DRUG INTERACTIONS

The following medications can prolong or potentiate the effects of anticoagulants and antagonize some of the therapeutic actions of Phytomenadione (vitamin K1): Phenylbutazone, Acetylsalicylic acid, Chloramphenicol, Sulfonamides (including sulfa / trimethoprim), Diazoxide, Allopurinol, Cimetidine, Metronidazole, Anabolic steroids, Erythromycin, Ketoconazole, Propanolol and thyroid drugs, therefore it is not recommended to use concomitantly with other pharmaceuticals.

CONTRAINDICATIONS

- Do not administer in animals with known hypersensitivity to Phytomenadione (Vitamin K1).
- Do not administer in patients with severe liver failure.
- Do not administer intravenously due to the risk of anaphylactic reactions.
- Do not administer intramuscularly due to the risk of bleeding and bruising.
- Do not administer the product for more than 4 administrations due to the risk of producing hemolysis.

PRECAUTIONS

Use during pregnancy, lactation and in breeding animals

Exogenous Phytomenadione (vitamin K1) enters the milk and crosses the placental barrier. There is no formal research on the safety of Phytomenadione during pregnancy, therefore, use it only according to the benefit / risk assessment made by the treating Veterinarian.

SPECIAL PRECAUTIONS FOR USE

Use during pregnancy, lactation and in breeding animals

Exogenous Phytomenadione (vitamin K1) enters the milk and crosses the placental barrier. There is no formal research on the safety of Phytomenadione during pregnancy, therefore, use it only according to the benefit / risk assessment made by the treating Veterinarian.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- In the case of ocular exposure, the eyes should be flushed with copious amounts of water at room temperature for at least 15 minutes. If irritation, pain, inflammation, tearing or photophobia persist, the patient should be evaluated by a doctor.
- In the case of dermal exposure, contaminated clothing should be removed and the exposed area washed thoroughly with soap and water. If there is irritation or pain the patient should be evaluated by a doctor.
- In the case of accidental injection, medical attention should be sought.

WARNINGS

Mantener alejado del alcance de los niños.

ADVERSE EFFECTS

Unwanted effects and adverse reactions:

High doses of Phytomenadione (vitamin K1) should be administered with caution due to the fact that Heinz body anemia has been reported in dogs that received doses of 4 mg / kg for 5 days.
In some dogs, cases of urticaria and abscess formation have been reported after subcutaneous administration of Phytomenadione (vitamin K1)

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Dispose of unused product remains in its original container, well closed. Dispose of the waste of this product with care together with household waste.

CONSERVATION

- Keep the product, closed or once opened, at a temperature between 2 ° C and 30 ° C, protected from light.
- Once opened, use the product within 2 weeks.
- Discard the unused product after that period of time.

CONDITION OF SALE

Sale with Veterinary Medical prescription.

PRESENTATION

20 mL vial

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 2329

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

CABATINA® ORAL SOLUTION

SOLUCIÓN ORAL.

ANTIHEMORRÁGICO PARA LA INTOXICACIÓN POR RODENTICIDAS ANTICOAGULANTES.

Technical Specification

SPECIES

Dogs and cats.

Cabatina® is a product indicated for the treatment of intoxications generated by anticoagulant rodenticides and hemorrhagic processes related to the defective formation of vitamin K-dependent coagulation factors.

DOSAGE FORM

Oral solution

THERAPEUTIC ACTION

Anti-haemorrhagic Anti-haemorrhagic for anticoagulant rodenticide poisoning.

COMPOSITION

Each 1 mL of product contains:
Vitamin K1 (Phytomenadione) ..... 20 mg
Excipients c.s.p 1mL

PROPERTIES

Vitamin K is a cofactor in the last stage of the hepatic synthesis of coagulation factors II (Prothrombin), VII (Proconvertin), IX (thromboplastin component of plasma) and X (Stuart factor). The process of formation of vitamin K-dependent clotting factors can be interrupted for various reasons, such as congenital diseases, vitamin K deficiency as a result of low dietary intake, malabsorption syndromes, broad-spectrum antibiotic therapies, or others.

In veterinary medicine, the most frequent cause of the cessation of K-dependent coagulation factor formation is intoxication by anticoagulant rodenticides. Vitamin K1 antagonizes the effects of anticoagulant rodenticidal agents and allows to reactivate the production process of K-dependent coagulation factors.

INDICATIONS

- Do not administer to animals with known hypersensitivity to vitamin K.
- Do not administer in animals with severe liver failure.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral.

Dosage:

Dose of the active principle:

Dogs: 2 mg / Kg of vitamin K1, once a day, for 3 weeks.
Cats: 5 mg / Kg of vitamin K1, once a day, for 3 weeks.

Product dosage:

Dogs: 1 mL of Cabatina® for every 10 Kg of body weight, once a day, for 3 weeks.
Cats: 1 mL Cabatina® for every 4 Kg of body weight, once a day, for 3 weeks.

Recommendations:
To improve its absorption, it is recommended to administer Cabatina® together with a tablespoon of a food rich in fatty acids, such as canned food for dogs or cats.

DRUG INTERACTIONS

The following medications can prolong or potentiate the effects of anticoagulants and antagonize some of the therapeutic actions of Vitamin K1: Phenylbutazone, Aspirin, Chloramphenicol, Sulfonamides (including Sulfa / Trimethoprim), Diazoxide, Allopurinol, Cimetidine, Metronidazole, anabolic steroids, Erythromycin , Ketoconazole, Propanolol and thyroid drugs.
The joint administration of Vaseline can decrease the absorption of Vitamin K.

CONTRAINDICATIONS

- Do not administer to animals with known hypersensitivity to vitamin K.
- Do not administer in animals with severe liver failure.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- In case of contact with the skin, it is recommended to wash hands with soap and plenty of water. If irritation develops and persists, see doctor.
- In the case of contact with the eyes, it is recommended to wash with plenty of water for 15 minutes. If irritation develops and persists, see doctor.
- In case of ingestion, do not induce vomiting. Get medical help.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Dispose of the waste of this product with care together with household waste. Contact the manufacturing company or companies specialized in waste disposal, to receive recommendations on the disposal of expired or unused products.

CONSERVATION

- Store at a temperature between 15 and 30 ° C, protected from light.
- Once opened, use within 8 weeks. Discard the unused product after that period of time.

CONDITION OF SALE

Sale with Veterinary Medical prescription.

PRESENTATION

25 mL bottle

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: SAG Reg. No. 2096
Peru: Reg. SENASA F.77.32.I.0053
Imported and distributed by Representaciones Durand SAC.
Av. Manuel Olguín N° 501 Office N° 604 Santiago de Surco Lima.
Bolivia: Reg. SENASAG PUV-No 009805/21
Imported and distributed by:
ZOO PHARMA VETERINARY INPUTS S.R.L.
Diaz Romero N° 1339 Miraflores Zone, La Paz-Bolivia
Telephone: 591 2223357
Costa Rica: Reg. No. MV-7129

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

CALFOMA® 12 - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

CALCIO, FOSFATO, MAGNESIO Y VITAMINA B12.

Technical Specification

SPECIES

Cattle, sheep, goats, pigs and horses.

Calfoma-12® is recommended in the syndrome of the fallen cow of metabolic origin: deficiencies of Calcium, Phosphorus, Magnesium and Vitamin B12. It is also indicated in lactation tetany, deficiency states or insufficient mineral intake, as well as for the toning of animals.

DOSAGE FORM

Injectable solution

THERAPEUTIC ACTION

Calcium, Phosphate, Magnesium and Vitamin B12.

COMPOSITION

Each 100 mL contains:
Calcium gluconate ..... 25.0 g
Sodium Glycerophosphate x 5.5 H2O ..... 1.0 g
Vitamin B12 ..... 3.5 mg
Magnesium Chloride x 6 H2O ..... 6.0 g
Excipients q.s.p ..... 100 mL

INDICATIONS

- Do not administer in dehydrated animals without prior hydration therapy.
- Do not administer in hyperexcited animals, due to the risk of cardiac arrest.

USE INSTRUCTIONS

Warm the container before administering the product. Inject aseptically. It should be administered by slow intravenous route, observing the reaction of the animal against any symptoms of tachycardia or arrhythmia.

ROUTE OD ADMINISTRATION AND DOSAGE

Route of administration: Endovenous, subcutaneous or intramuscular.

Dosage:

- Bovines: 250 to 800 mL in hypocalcemia.
- Calves between 50 to 100 Kg: 25 to 60 mL for toning.
- Sheep and goats: 25 to 60 mL in hypocalcemia, hypomagnesemia and / or toning.
- Pigs: 25 to 60 mL for toning.
- Piglets between 10 to 30 Kg: 5 to 10 mL for toning.
- Horses: 250 to 800 mL for toning.

DRUG INTERACTIONS

- Calcium prolongs or enhances the effects of Tubocurarine.
- Animals receiving parenteral calcium and potassium supplements are at higher risk of cardiac arrhythmias: administer with care.
- Excessive intake of vitamin A can stimulate skeletal calcium depletion and cause hypercalcemia.
- In concurrent use of high doses of vitamin D or its analogues it can increase the absorption of Calcium and promote hypercalcemia.

CONTRAINDICATIONS

- Do not administer in dehydrated animals without prior hydration therapy.
- Do not administer in hyperexcited animals, due to the risk of cardiac arrest.

PRECAUTIONS

Special warnings and precautions for use:

- Administration must be supervised by a Veterinarian.
- When the route of administration chosen is intravenous, rapid administration should be avoided, as it can cause ventricular fibrillation, the risk of injectable Calcium.
- Due to its high Calcium content, Calfoma-12® can crystallize at low temperatures. In this case, heat the container until the crystals formed dissolve.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR USE

Special warnings and precautions for use:

- Administration must be supervised by a Veterinarian.
- When the route of administration chosen is intravenous, rapid administration should be avoided, as it can cause ventricular fibrillation, the risk of injectable Calcium.
- Due to its high Calcium content, Calfoma-12® can crystallize at low temperatures. In this case, heat the container until the crystals formed dissolve.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- In the case of ocular exposure, irrigate the eyes with abundant water. If irritation exists and persists, consult a doctor.
- In the case of dermal exposure, contaminated clothing should be removed and the exposed area washed with water. If irritation exists and it persists, consult a doctor.
- In the case of accidental injection, seek medical attention.

ADVERSE EFFECTS

Inflammation may be seen at the site of the subcutaneous or intramuscular injection, which gradually subsides.

GUARD PERIOD

0 days.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory

CONSERVATION

Store in a cool and dry place, at room temperature between 15º and 30ºC.

CONDITION OF SALE

Sale with Veterinary Medical prescription.

PRESENTATION

100 mL vial and 500 mL serum bottle

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

- Chile: Reg. SAG N° 587
- El Salvador: Reg. Nº VET.2003-03-2696

COUNTRIES WHERE IT IS MARKETED

Distribution in El Salvador:
Rafael Alfredo Alfaro Castillo.
8th C. Pte. Pje. Moreno N ° 112, Col. Flor Blanca.
San Salvador, El Salvador.

CALFOMA® PLUS - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

CALCIO, FÓSFORO, MAGNESIO Y DEXTROSA CON VITAMINA B12.

Technical Specification

SPECIES

Bovines.

It is recommended in the syndrome of the fallen cow of metabolic origin, deficiencies of Calcium, Phosphorus, Magnesium, Dextrose and Vitamin B12. It is also indicated in tetany of lactation and deficiency states or deficiencies in the contribution of minerals.

DOSAGE FORM

Injectable Solution

THERAPEUTIC ACTION

Calcium, Phosphorus, Magnesium and Dextrose with Vitamin B12

COMPOSITION

Each 100 mL of solution for injection contains:
Calcium Gluconate Monohydrate ..... 25.0 g
Sodium Hypophosphite Monohydrate ..... 0.12 g
Magnesium Chloride Hexahydrate ..... 6.0 g
Sodium Glycerophosphate x 5.5 H2O ..... 1.0 g
Dextrose..... 5.0 g
Vitamin B12 ..... 3.5 mg
Excipients q.s.p ..... 100 mL

INDICATIONS

- Do not administer in dehydrated animals without prior hydration therapy.
- Do not administer in hyperexcited animals due to the risk of cardiac arrest.

USE INSTRUCTIONS

Warm the container before administering the product. Inject aseptically. It should be administered by slow intravenous route, observing the reaction of the animal against any symptoms of tachycardia or arrhythmia.

ROUTE OD ADMINISTRATION AND DOSAGE

Route of administration: slow intravenous

Product dosage:
Administer between 500 to 1,000 mL of Calfoma® Plus per animal.

CONTRAINDICATIONS

- Do not administer in dehydrated animals without prior hydration therapy.
- Do not administer in hyperexcited animals due to the risk of cardiac arrest.

PRECAUTIONS

- Do not administer by rapid intravenous route, due to the potential danger of ventricular systole.
- It is recommended to administer in combination with a cardiac tonic.

SPECIAL PRECAUTIONS FOR USE

- Do not administer by rapid intravenous route, due to the potential danger of ventricular systole.
- It is recommended to administer in combination with a cardiac tonic.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a cool and dry place, at room temperature between 15º and 30ºC.

CONDITION OF SALE

Veterinary Medical prescription.

PRESENTATION

500 mL vial

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

- Chile: Reg. SAG N° 1384
- Costa Rica: Reg. N° MAG CL4-32-02-3647
- Rep. Dominicana: Reg. Nº 5606

CALMEDRAG® ORAL TABLETS

COMPRIMIDO ORAL.

ANSIOLÍTICO.

Technical Specification

SPECIES

Dogs.

Recommended as a treatment for the separation anxiety.
The treatment with Calmedrag® must be supported with conduct management therapy.

DOSAGE FORM

Oral tablets.

THERAPEUTIC ACTION

Anxiolytic.

COMPOSITION

Each tablet contains:
Clomipramine Hydrochloride .....20 mg
(Equivalent to 17.9 mg of Clomipramine base)
Excipients q.s.p.....1 tablet.

PROPERTIES

Clomipramine Hydrochloride is a tricyclic anti-depressant that inhibits the reuptake of presynaptic serotonin and norepinephrine and with this produces anxiolytic, anti-compulsive, anti-aggressive and anti-depressant effects. As a result can be seen significant changes in the behavior of the dogs subjected to therapies with Clomipramine.

INDICATIONS

- Do not administer to animals with known hypersensitivity to tricyclic agents.
- Do not administer to animals weighing less than 2.5 Kg.
- Do not administer to male breeding dogs.
- Do not administer to pregnant or nursing females.
- Do not administer to puppies less than 6 month old.
- Do not administer concomitantly with MAO inhibitors (Selegiline, Amitraz)

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route of administration.

Place the tablet as far as possible on the muzzle of the dog, keep the muzzle closed and stimulate the swallowing. It is recommended at the beginning of the therapy, to administer the product along with the food to reduce the possibility of side effects such as vomiting.

Active ingredient dose:

- 2 mg of Clomipramine Hydrochloride per 10 kg of body weight (2 mg/Kg) every 12 hours for 2-3 months.

Product dose:

- 1 tablet of CALMEDRAG® per each 10 Kg of body weight (1 tablet/10 Kg) every 12 hours for 2-3 months.
  It is advisable to suspend therapy gradually.

OVERDOSE: Clomipramine in general is a fairy safe medication and well tolerated by dogs. In this species the lethal dose is approximately between 50 and 100 mg/Kg per day, that is to say 12.5 to 25 times the recommended therapeutic dose.

Notwithstanding the foregoing, the overdose with tricyclic antidepressant can be life threatening (arrhythmia, convulsions, cardiac and respiratory arrest).

There is no known antidote for Clomipramine.

DRUG INTERACTIONS

- Due to the additive effects, Clomipramine must be administered with caution when used concomitantly with other anticholinergic agents or CNS depressants.
- Tricyclic antidepressants used with anti-thyroid agents may increase the potential risk of agranulocytosis.
- The cimetidine may inhibit the metabolism of the tricyclic antidepressants and increase the risk of toxicity.
- Its use in combination with sympathomimetic agents may increase the risk of cardiac effects (arrhythmia, hypertension, hyperpyrexia).
- Do not administer concomitantly with MAO inhibitors (Selegiline, Amitraz)

CONTRAINDICATIONS

- Do not administer to animals with known hypersensitivity to tricyclic agents.
- Do not administer to animals weighing less than 2.5 Kg.
- Do not administer to male breeding dogs.
- Do not administer to pregnant or nursing females.
- Do not administer to puppies less than 6 month old.
- Do not administer concomitantly with MAO inhibitors (Selegiline, Amitraz)

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Wash hands thoroughly after handling and/or administering this product.
- Keep out of reach of children.
- The accidental ingestion should be considered dangerous.
- People with known sensitivity to Clomipramine must exercise caution handling this product.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

- The symptoms and adverse signs more frequently observed include nausea, vomiting and lethargy or transient drowsiness.
- The sedation typically occurs at the beginning of the therapy and is usually self-limiting, as soon as the dog becomes tolerant to it.
- The vomiting has been reported specially in dogs that have received the Clomipramine in an empty stomach, therefore it is recommended that administer it along with the food.
- There could be, although unlikely, anorexia, diarrhea, hyperactivity of liver enzymes and some cholinergic effects (for example, dry mouth).

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
Empty containers can be discarded as household waste, without any special precautions. Do not dispose of containers with product residues on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store in a cool and dry place, at room temperature between 15º y 30°C.

CONDITION OF SALE

Chile:Venta bajo receta médico veterinaria retenida
Uruguay: venta exclusiva bajo receta msp (receta verde)

PRESENTATION

Case with 30 tablets.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 2178
Uruguay: Reg. MGAP N° 2017A00466

COUNTRIES WHERE IT IS MARKETED

Importer in Uruguay by:

VIVAFIL S.A.
RIO NEGRO 1107 Montevideo - Uruguay, TEL 29001112
grupotecnovet@gmail.com
Technical Director: DMTV Diego Cuadrado.

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

CALVIPET® ORAL POWDER

POLVO ORAL.

CALCIO VITAMINADO EN POLVO PARA PERROS.

Technical Specification

SPECIES

Dogs.

Calvipet® is a food supplement of calcium, Phosphorus and Vitamin D, which is specially recommended for the prevention and adjustment of dietary deficiencies in dogs.

DOSAGE FORM

Oral powder.

THERAPEUTIC ACTION

Calcium vitaminic powder for dogs.

COMPOSITION

Each 100 g contains:
Calcium Phosphate Dibasic Dihydrate.....14 g
Calcium Lactate Pentahydrate.....10 g
Vitamin D3.....0.2 g (20.000 IU)
Excipients q.s.p.....100 g

INDICATIONS

- Do not administer in dogs in maintenance that feed on balanced diets in Calcium, Phosphorus or Vitamin D.
- Do not administer in cases of renal failure, renal lithiasis or hypercalcemia.
- Do not administer in cases of secondary hyperparathyroidism.

ROUTE OD ADMINISTRATION AND DOSAGE

Route of administration: Oral route, mixed with food.

Product dosage: (1 teaspoon = 5 g)
a) Prevention of dietary deficiencies: 1 to 2 tsp. / 10 Kg once a day for 1 month.
b) Correction of dietary deficiencies: 2 to 4 tsp. / 10 kg once a day for 3 months.

CONTRAINDICATIONS

- Do not administer in dogs in maintenance that feed on balanced diets in Calcium, Phosphorus or Vitamin D.
- Do not administer in cases of renal failure, renal lithiasis or hypercalcemia.
- Do not administer in cases of secondary hyperparathyroidism.

PRECAUTIONS

- This product should not be administered indefinitely.
- Oral Tetracycline treatments should be administered at least 1 to 2 hours before or after administering the product.

SPECIAL PRECAUTIONS FOR USE

- This product should not be administered indefinitely.
- Oral Tetracycline treatments should be administered at least 1 to 2 hours before or after administering the product.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions.
Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 ° C, protected from light.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

Pot with 100 g

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg SAG No. 958

CANIFORT® ORAL TABLETS

COMPRIMIDO ORAL.

ANTIPARASITARIO INTERNO DE AMPLIO ESPECTRO PARA PERROS.

Technical Specification

SPECIES

Dogs.

Internal broad-spectrum antiparasitic, effective against Toxocara canis, *Toxascaris leonina, Uncinaria stenocephala, Ancylostoma caninum, Trichuris vulpis, Taenia hydatigena, Taenia pisiformis, Dipylidium caninum and Giardia spp.*

DOSAGE FORM

Oral tablet.

THERAPEUTIC ACTION

Internal broad spectrum dewormer for dogs.

COMPOSITION

Each tablet contains:
Praziquantel ..... 50 mg
Pirantel Pamoate ..... 144 mg
(Equivalent to 50 mg of Pirantel)
Febantel ..... 150 mg
Excipients q.s.p ..... 1 tablet

INDICATIONS

- Do not administer to dogs younger than 4 weeks.
- Do not administer to pregnant females.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration.

Dosage of the active ingredients:
Febantel 15 mg / Kg; Pirantel 5 mg / Kg; Praziquantel 5 mg / Kg of weight, in a single dose.
Product dosage:
1 tablet for every 10 Kg of weight, in a single dose. For the treatment of giardasis, administer one tablet for every 10 kg of weight for three consecutive days.

CONTRAINDICATIONS

- Do not administer to dogs younger than 4 weeks.
- Do not administer to pregnant females.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

Vomiting may occur in dogs treated with Pirantel and / or Febantel, although it is highly unlikely.
Oral use, Praziquantel can cause anorexia, vomiting, lethargy or diarrhea in dogs, but the incidence of these signs is less than 5%.

OBSERVATIONS

Special precautions for the disposal of waste material:
Discard the remains of unused product in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of water. Dispose of this product with caution with household waste.

CONSERVATION

Store at a temperature between 2º and 30ºC, sheltered from light, in a cool and dry place.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Box with 1 or 50 tablets.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 0349
- Costa Rica: Reg. No. MAG CL4-42-10-3630
- El Salvador: Reg. No. 2003-03-2682
- Panama: Reg. No. RF-4111-018

COUNTRIES WHERE IT IS MARKETED

Distribution in El Salvador:
Rafael Alfredo Alfaro Castillo.
8a C. Pte. Pje. Moreno N ° 112, Col. Flor Blanca.
San Salvador, El Salvador.

CANISH® BALSAMIC SHAMPOO - SHAMPOO AND CONDITIONER

SHAMPOO Y ACONDICIONADOR.

DOS EN UNO, LIMPIA Y ACONDICIONA EL PELAJE.

Technical Specification

SPECIES

Dogs.

Restores and maintains the natural silkiness and moisture of the pet's skin and coat.
Balanced according to the pH of the skin. CANISH® Balsamic is hypoallergenic and does not contain irritating chemicals.

DOSAGE FORM

Balsamic Shampoo and Conditioner

THERAPEUTIC ACTION

Two in one, cleans and conditions the coat

INGREDIENTS

Purified Water, Sodium Lauryl Ether Sulfate 70, Hydrolyzed Collagen, Hydroxypropyl Lauryl Diammonium, Cocamide DEA, Glycol Distearate / Laureth-4 / Cocamidopropyl Betaine, Sodium Chloride, Disodium EDTA, Authorized Fragrances and Preservatives.

PROPERTIES

Shampoo and Conditioner

- Clean and detangle in one go
- Easy to style
- Smoothness
- Intense shine

USE INSTRUCTIONS

- Wet your pet with warm water and pour sufficient quantity of CANISH® Balsamic Shampoo on the back of the animal.
- Form abundant lather, massaging gently.
- Wash using a plentiful quantity of water.
- Apply again CANISH® Balsamic Shampoo, if necessary.
- Rinse with water until the lather disappears.

CONSERVATION

Store in a cool and dry place, out of direct light, below 30°C

PRESENTATION

390 mL bottle

PREPARED BY

Drag Pharma Laboratory

RECORDS

Rep. Dominicana: Reg. Nº 8348
Costa Rica: Reg. MAG Nº CL-4-45-5-6012
Bolivia: Reg. SENASAG PUV N° 008262/18

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

CANISH® DRY FOAM - SHAMPOO

SHAMPOO EN ESPUMA SECA.

LIMPIEZA EN SECO PARA PERROS.

Technical Specification

SPECIES

Dogs and cats.

DOSAGE FORM

Dry Foam Shampoo

THERAPEUTIC ACTION

no rinse foaming dry shampoo for Dogs

INGREDIENTS

Purified Water, Sodium Lauryl Ether Sulfate, Cocamidopropyl Betaine, Propylene Glycol, PEG-12 Dimethicone, Sodium Chloride, Disodium EDTA, Authorized Fragrances and Preservatives.

PROPERTIES

Cleanness, gloss and softness for your pet coat, without needing to rinse.
Wild Aroma

USE INSTRUCTIONS

- Shake the bottle vigorously, then spread small amounts of foam over the coat of the animal.
- Rub it well with a clean, dry towel.
- Finally brush your pet's hair to get a neat coat and regain its shine.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

- For external use.

CONSERVATION

- Store in a cool and dry place, out of direct light, below 30°C.

PRESENTATION

160 g

PREPARED BY

Drag Pharma Laboratory

RECORDS

Rep. Dominicana: Reg. Nº 8347
Costa Rica: Reg. MAG Nº CL-4-45-14-6019
Bolivia: Reg. SENASAG PUV N° 008265/18

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

CANISH® EXTRA GLOSS - SHAMPOO

SHAMPOO.

SHAMPOO PARA PERROS.

Technical Specification

SPECIES

Dogs.

CANISH® Extra Shine Shampoo has been specially formulated to leave a protective film on your dog's coat after washing, enhancing its natural color and giving it softness and intense shine.
This protective action, together with the effect of moisturizing agents and the balanced pH of the formula, allows to maintain a clean coat and a lasting luster for longer, making your pet look healthier and more vital. Its soft essence of red fruits will leave a pleasant aroma on your pet.

DOSAGE FORM

Shampoo.

THERAPEUTIC ACTION

Dog Shampoo.

INGREDIENTS

Purified Water, Sodium Lauryl Ether Sulfate, Cocamide DEA, Sodium Chloride, PEG-12 Dimethicone, Authorized Fragrance, Disodium EDTA, Isothiazolinones, Anhydrous Citric Acid, FD&C Yellow Color No. 5.

PROPERTIES

EXTRA GLOSS

- Intense shine
- Long-lasting protection against dust and dirt
- Smoothness
- Moistening

USE INSTRUCTIONS

- Wet your pet with warm water and pour sufficient quantity of CANISH® Extra Gloss on the back.
- Form abundant lather, massaging gently.
- Wash using a plentiful quantity of water.
- Apply again CANISH® Extra Gloss, if necessary.
- Rinse with water until the lather disappears.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a cool and dry place, out of direct light, below 30°C

PRESENTATION

390 mL bottle

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Rep. Dominicana: Reg. Nº 8350
Bolivia: Reg. SENASAG PUV N° 008273/18

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

CANISH® HERBAL EXTRACT - SHAMPOO

SHAMPOO.

SHAMPOO PARA PERROS.

Technical Specification

SPECIES

Dogs.

CANISH® Herbal Extract Shampoo is a formula enriched with herbal extracts such as Salvia, Rosemary and Nettle, which provide a natural toning effect to your pet's coat. Each herbal component has a stimulating action on the hair follicles, strengthening hair growth from the roots. Rosemary also helps to eliminate dryness and has an excellent conditioning effect; Sage reduces excess fat from the dog's coat and Nettle provides softness and shine.

DOSAGE FORM

Shampoo

THERAPEUTIC ACTION

Dog shampoo.

INGREDIENTS

Purified Water, Sodium Lauryl Ether Sulfate, Cocamide DEA, Extracts of sage leaves (Salvia officinalis), extract of nettle (Urtica dioica), extract of rosemary leaf (Rosmarinus officinalis) and extract of bell pepper (Capsicum annuum); Sodium chloride; Disodium EDTA; Authorized Colors, Fragrances and Preservatives.

PROPERTIES

HERBAL EXTRACT

- Invigorating effect on the coat.
- Rosemary, eliminates dryness, excellent conditioner.
- Salvia, reduces excess fat.
- Nettle, strengthens growth.
- Not irritating.

USE INSTRUCTIONS

- Wet your pet with warm water and pour sufficient quantity of CANISH® Herbal Extracts on the back.
- Form abundant lather, massaging gently.
- Wash using a plentiful quantity of water.
- Apply again CANISH® Herbal Extracts, if necessary.
- Rinse with water until the lather disappears.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a cool and dry place, out of direct light, below 30°C

PRESENTATION

390 mL bottle

PREPARED BY

Drag Pharma Laboratory

RECORDS

Rep. Dominicana: Reg. Nº 8346
Bolivia: Reg. SENASAG PUV N° 008260/18

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

CANISH® HYPOALLERGENIC - SHAMPOO

SHAMPOO.

SHAMPOO PARA PERROS.

Technical Specification

SPECIES

Dogs.

Its hypoallergenic components and its balanced pH make it suitable for bathing all those races with sensitive skin or with recurrent dermatological disorders.

DOSAGE FORM

Shampoo

THERAPEUTIC ACTION

Dog shampoo

INGREDIENTS

Purified Water, Sodium Lauryl Ether Sulfate 70, Cocamide DEA, Sodium Chloride, Fragrances and Authorized preservatives.

PROPERTIES

Hypoallergenic Shampoo

- Frequent use
- balanced pH
- Smoothness

USE INSTRUCTIONS

- Wet your pet with warm water and pour sufficient quantity of CANISH® Hypoallergenic on the back.
- Form abundant lather, massaging gently.
- Wash using a plentiful quantity of water.
- Apply again CANISH® Hypoallergenic, if necessary.
- Rinse with water until the lather disappears.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a cool and dry place, out of direct light, below 30°C

PRESENTATION

390 mL bottle

PREPARED BY

Drag Pharma Laboratory

RECORDS

Costa Rica: Reg. MAG CL-4-45-4-6238
Rep. Dominicana: Reg. Nº 8349
Bolivia: Reg. SENASAG PUV N° 008261/18

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

CAN-OUT® SPRAY SOLUTION

SOLUCIÓN SPRAY.

SOLUCIÓN SPRAY AMARGANTE ANTIMORDISQUEOS DE PERROS. AYUDA A CORREGIR LOS MALOS HÁBITOS DE SU MASCOTA.

Technical Specification

SPECIES

DOSAGE FORM

Spray solution.

THERAPEUTIC ACTION

Bitter spray solution anti-chew for dogs. Help to change the bad habits of your pet.

COMPOSITION

Each 100 mL of product contains:
Denatonium Benzoate.....60 mg
Excipients q.s.p.....100 mL

PROPERTIES

Can-Out® is a bitter spray solution with repellent action against chewing of dogs, which is applied on surfaces such as furniture, curtains, shoes, clothes, toys and others, for preventing destruction caused by dog's bites, specially puppies.

Can-Out® is also effective when used on dressings or patches in the animal, avoiding to use an Elizabethan collar.

Can-Out® does not leave residues or stains on the surfaces treated.

Can-Out® contains alcohol; therefore, it should not be used in open wounds.

Can-Out® contain Denatonium Benzoate, a bitter substance that causes an immediate rejection in the animal when the impregnated surface is chewed.

The proper use of this product does not pose any risk to your pet.

USE INSTRUCTIONS

Apply the spray on the object or surface that you want the dog not to bite. Apply an amount enough in order to completely impregnate the object. Thus, when the dog approaches to bite, will reject. Apply at first several times a day and then once a day.

WARNINGS

- Producto inflamable.
- Mantener fuera del alcance de los niños.
- No aplicar directamente a su mascota.
- Probar el producto en un sector oculto para verificar la resistencia de los colores.

CONSERVATION

- Store in a cool, dry place, protected from light, at no more than 30ºC.
- Keep away from heat sources.

PRESENTATION

100 mL bottle with spray bottle.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Costa Rica: Reg. MAG CL-4-13-1-6020

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Costa Rica by:
Proventas de Cartago S.R.L. 100 meters east Hogares Crea, San Blas. Carthage.

CAVIVET® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

TÓNICO RECONSTITUYENTE.

Technical Specification

SPECIES

Bovines, horses, pigs and sheep.

Tonic restorative adjuvant in the treatment of various pathologies.

- **Horses:** Failures of development and growth; fertility problems; improvement of skin and coat, improvement in the training and in preparation for rodeo participation.
- **Bovine:** Strengthening of calves, strengthening of cows in the last trimester of gestation; fertility problems; performance failure in bulls; and treatment of milk fever in cattle.
- **Pigs:** Strengthening of piglets, breeding sows and boars in mounting.
- **Ovine:** Strengthening of rams and sheep in the last trimester of gestation.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Restorative tonic

COMPOSITION

Each 100 mL of injectable solution contains:
Calcium Gluconate Monohydrate.....5.0000 g
Calcium D-Sacarate Tetrahydrate.....0.0800 g
Calcium Levulinate Dihydrate.....1.0000 g
Calcium Glycerophosphate Anhydrous.....0.5000 g
Magnesium Glycerophosphate Dihydrate.....1.5000 g
Sodium Glycerophosphate x 5.5 H2O.....0.8000 g
Sodium selenite Pentahydrate.....0.0333 g
Cobalt Chloride Hexahydrate.....0.0800 g
Potassium Chloride.....0.4000 g
Excipients q.s.p.....100 mL

INDICATIONS

Do not administer simultaneously with Tetracyclines.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration route:
Intramuscular, subcutaneous, slow intravenous.

Dose:

- **Cattle and horses:** 15 - 20 mL in a single dose. The treatment can be repeated after one week.
- **Sheep and pigs:** 10 mL in a single dose. The treatment can be repeated after one week.

CONTRAINDICATIONS

Do not administer simultaneously with Tetracyclines.

WARNINGS

Advertencias y precauciones especiales de uso:

- La administración endovenosa podría provocar problemas de flebitis.
- La administración endovenosa debe hacerse de forma lenta por riesgo de bradicardia o paro cardíaco.
- Mantener fuera del alcance de los niños y animales domésticos.

GUARD PERIOD

Meat: 0 days.
Milk: 0 days.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
Empty containers can be discarded as household waste, without any special precautions.
Do not dispose of containers with product residues on the ground or water courses.
For expired or unused products contact the manufacturer laboratory.

CONSERVATION

Store between 15 ° and 30 ° C, protected from light.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

250 mL ampoule bottle

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 937
- Costa Rica: Reg. No. MAG CL4-32-02-3648

CLINDABONE® ORAL TABLETS

COMPRIMIDO ORAL.

ANTIBIÓTICO.

Technical Specification

SPECIES

Dogs and cats.

Recommended for the treatment of infections caused by bacteria sensitive to Clindamycin, such as *Staphylococcus*, *Streptococcus*, and anaerobic bacteria as *Bacteroides spp.*, *Fusobacterium spp.* , *Peptoestreptococcus spp.* , *Clostridium perfringens* and many species of *Propionibacterium spp.* It also presents activity against *Toxoplasma gondii*. It use is recommended for the treatment of infections caused by bacteria sensitive to Clindamycin in dogs and cats. In dogs, it is recommended as elective therapy for oral infections, pyoderma and osteomyelitis. In cats, it is recommended especially for oral infection treatments. Clindamycin is also recommended in cases of toxoplasmosis in dogs and cats, since allows the improvement of the clinical signs associated with the active infection with *Toxoplasma gondii*.

DOSAGE FORM

Oral tablets.

THERAPEUTIC ACTION

Antibiotic.

COMPOSITION

Each Tablet of 620 mg contains:
Clindamycin Hydrochloride .....195.04 mg
(Equivalent to 165 mg of basic Clindamycin)
Excipients q.s.p.....620 mg

PROPERTIES

Clindamycin is an antibiotic belonging to the Lincosamide group, which corresponds to a semi-synthetic modification of Lincomycin. It has a spectrum and action similar to Erythromycin, that is, gram positive bacteria, especially staphylococci, streptococci and anaerobic bacteria. Lincosamides are useful for the treatment of infections resistant to Penicillin, Erythromycin and Cephalosporins. Clindamycin inhibits bacterial protein synthesis, altering translocation. It binds to the 50S subunit of the bacterial ribosome, preventing translocation of transfer RNA from site A to site P of the ribosome, preventing the formation of peptide bonds and generating the inhibition of the protein synthesis of the microorganism.

INDICATIONS

- Do not administer in animals with hypersensitivity to Lincomycin and Clindamycin.
- Do not use in newborn puppies or kittens.
- Do not administer to cats weighing less than 2 kg.
- Do not administer concomitantly with other antibiotics such as Chloramphenicol or Macrolides.
- Do not use in other species than those indicated since their intake can cause serious gastrointestinal disorders.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration.

Active ingredient dose:

The recommended dose for dogs and cats for the treatment of **oral infections** is: 11 mg/Kg of weight every 24 hours or 5.5 mg/Kg of weight every 12 hours, for 5 days. For the treatment of **osteomyelitis** in dogs is recommended a dose of 11 mg/Kg of weight every 12 hours, for 28 days. The beginning active dose for the treatment of **pyoderma** in dogs is 11mg/Kg every 24 hours or 5.5 mg/Kg of weight every 12 hours, for a minimum of 21 consecutive days or at least 7 days after the apparent clinical cure. For the treatment of **toxoplasmosis** is recommended a dose of Clindamycin for dogs and cats of 10 to 40 mg/Kg and 25 to 50 mg/Kg, respectively, divided into 2 daily doses, for 14 to 21 days.

Product dose:

Oral infections (Periodontitis):

- Dogs: ½ Tablet for every 15 Kg of weight twice a day, or 1 tablet for every 15 Kg of weight, once a day for 5 days
- Cats: ¼ Tablet for every 4 kg of weight, once a day for 5 days.

Osteomyelitis:

- Dogs: 1 tablet for every 15 Kg of weight, twice a day for 28 days.

Pyoderma:

- Dogs: ½ Tablet for every 15 Kg of weight, twice a day or 1 tablet for every 15 Kg of weight, once a day for 21 consecutive days or at least 7 days after the apparent clinical cure.

Toxoplasmosis:

- Dogs: 1 tablet for every 10 Kg of weight, twice a day, for 14 to 21 days.
- Cats: ¼ Tablet for every 2 to 3 Kg of weight, twice a day, for 14 to 21 days.

DRUG INTERACTIONS

- Lincosamides have a neuromuscular additive effect, when applied with anesthetic agents and relaxants of the skeletal musculature.
- The association with Kaolin-Pectin does not allow absorption from the gastrointestinal tract.
- It should not be combined with other bactericidal agents, nor with macrolides, since there may be antagonisms or nephrotoxicity.
- Cyclosporins: Clindamycin can reduce levels.
- Erythromycin: antagonism in vitro when administered with Clindamycin.

CONTRAINDICATIONS

- Do not administer in animals with hypersensitivity to Lincomycin and Clindamycin.
- Do not use in newborn puppies or kittens.
- Do not administer to cats weighing less than 2 kg.
- Do not administer concomitantly with other antibiotics such as Chloramphenicol or Macrolides.
- Do not use in other species than those indicated since their intake can cause serious gastrointestinal disorders.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Wash hands after administering the product.
- In case of accidental ingestion, do not induce vomiting, it is recommended to seek immediate medical attention.
- In case of contact with the eyes, wash with plenty of water for at least 15 minutes.
- In the case of any type of irritation, it is recommended to go to the doctor.
- If there is any type of skin irritation, it is recommended to go to the doctor.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

Prolonged use of Clindamycin may occasionally cause anorexia, vomiting, and diarrhea.

OBSERVATIONS

Use during pregnancy and lactation:
Do not administer to pregnant or lactating females.

Special precautions for the disposal of unused product or waste material:
Discard the remains of unused product in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of natural water. Dispose of this product with caution with household waste.

CONSERVATION

Store at a temperature between 15 and 30 ° C, protected from light.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Box containing 20 tablets.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 1959-B
Uruguay: Reg. MGAP Nº 2018A00613
Perú: Reg. SENASA F.82.21.I.0448

COUNTRIES WHERE IT IS MARKETED

Importer in Uruguay by:

VIVAFIL S.A.
RIO NEGRO 1107 Montevideo
Uruguay, TEL 29001112
grupotecnovet@gmail.com
Director Técnico: DMTV Diego Cuadrado.

Imported and Distributed in Peru by:

Representaciones Durand SAC.
Av. Manuel Olguín N° 501 Oficina N° 604
Santiago de Surco Lima.

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

COLIMIC® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ESPASMOLÍTICO.

Technical Specification

SPECIES

Cattle, horses, pigs and sheep.

Antispasmodic, relaxant of the smooth muscles in states of gastrointestinal colic and as an adjunct in diarrhea and vomiting.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Spasmolytic.

COMPOSITION

Each 10 mL of solution contains:
Papaverine Hydrochloride ..... 300 mg
(Equivalent to 271 mg of Papaverine)
Atropine Sulfate ..... 30 mg
(Equivalent to 26 mg of Atropine)
Excipients q.s.p ..... 10 mL

INDICATIONS

- Do not administer in animals with glaucoma and heart failure.
- Do not administer in cases of intestinal obstruction.
- Do not administer in animals with known hypersensitivity to Atropine or Papaverine.
- Do not administer during pregnancy or while breastfeeding.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration routes:
Intramuscular or subcutaneous administration

Active ingredients dose:

Atropine:

- Bovine and horses: 0.03 – 0.06 mg/Kg
- Pigs: 0.1 mg/Kg
- Ovine: 0.03-0.3 mg/Kg

Papaverine:

- Bovine and horses: 0.3-0.54 mg/Kg
- Pigs: 0.11 mg/Kg
- Ovine: 0.3 – 3.3 mg/Kg

Product dose:

- Bovine and horses: 1-2 mL/100 Kg of weight, in a single dose.
- Pigs: 4 mL / 100 Kg of weight, in a single dose.
- Ovine: 1-12 mL/100 Kg of weight, in a single dose.

DRUG INTERACTIONS

- The following actives can increase the activity or toxicity of Atropine and its derivatives: Amantadine, other anticholinergic agents, anticholinergic muscle relaxants, antihistamines (eg Diphenhydramine), Disopyramide, Meperidine, Phenothiaripzines, Procainamide, Primidone, Tricyclic antidepressants (eg Amiticiline antidepressants (eg Amiticiline antidepressants) , Clomipramine).
- The use of Atropine with alpha-2 agonists (eg Dexmedetomidine, Medetomidine) can significantly increase blood pressure, heart rate and the incidence of arrhythmias.
- Atropine can aggravate some symptoms caused by Amitraz intoxication; leading to hypertension and increased inhibition of peristalsis.
- Long-term use of Atropine in conjunction with corticosteroids can increase intraocular pressure.
- Atropine and its derivatives can antagonize the actions of metoclopramide.

CONTRAINDICATIONS

- Do not administer in animals with glaucoma and heart failure.
- Do not administer in cases of intestinal obstruction.
- Do not administer in animals with known hypersensitivity to Atropine or Papaverine.
- Do not administer during pregnancy or while breastfeeding.

PRECAUTIONS

- Anticholinergic agents should be administered with caution in geriatric animals, with kidney and liver disease or those with esophageal reflux.
- Administer with caution in horses as it can induce colic.

SPECIAL PRECAUTIONS FOR USE

- Anticholinergic agents should be administered with caution in geriatric animals, with kidney and liver disease or those with esophageal reflux.
- Administer with caution in horses as it can induce colic.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

Mydriasis, dry mucous membranes and intestinal atony in variable degrees. The most frequently described adverse effects are ruminal bloating in ruminants and tachycardia in horses.

GUARD PERIOD

Meat and milk: 0 days.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Dispose of this product waste carefully with household waste. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 °C, protected from light.
Once the container is opened, use within 28 days. Discard unused product after that time frame.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

10 mL and 50 mL vial

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No. 928

COLONIA ANIMAL HEALTH® BLUE

COLONIA PARA PERROS.

Technical Specification

SPECIES

Dogs

THERAPEUTIC ACTION

Cologne for dogs

INGREDIENTS

Demineralized water, Dimethicone PEG 12, Hydrogenated castor oil PEG 40, fragrances and authorized preservatives.

PROPERTIES

- Fragrance specially formulated for dogs from 2 months of age.
- It does not alter the smell of the animal.
- With soft touches of wood and natural forest.

USE INSTRUCTIONS

Apply sufficient quantity directly on the back and / or clothes of the pet.
Ideal to be used after the usual bath.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

FOR EXTERNAL USE ONLY

CONSERVATION

Store in a cool and dry place, protected from light, at no more than 30ºC.

PRESENTATION

180 mL spray bottle

COLONIA ANIMAL HEALTH® PUPPY

COLONIA PARA PERROS.

Technical Specification

SPECIES

Dogs

THERAPEUTIC ACTION

Cologne for dogs

INGREDIENTS

Demineralized water, Dimethicone PEG 12, Hydrogenated castor oil PEG 40, fragrances and authorized preservatives.

PROPERTIES

- Fragrance specially formulated for dogs from 2 months of age.
- It does not alter the smell of the animal.
- With natural floral chords and hints of enveloping warmth.

USE INSTRUCTIONS

Apply sufficient quantity directly on the back and / or clothes of the pet.
Ideal to be used after the usual bath.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

FOR EXTERNAL USE ONLY

CONSERVATION

Store in a cool and dry place, protected from light, at no more than 30ºC.

PRESENTATION

180 mL spray bottle

COLONIA ANIMAL HEALTH® VIOLET

COLONIA PARA PERROS.

Technical Specification

SPECIES

Dogs.

THERAPEUTIC ACTION

Cologne for dogs

INGREDIENTS

Demineralized water, Dimethicone PEG 12, Hydrogenated castor oil PEG 40, fragrances and authorized preservatives.

PROPERTIES

- Fragrance specially formulated for dogs from 2 months of age.
- It does not alter the smell of the animal.
- With the magic of a sweet aroma of melon and berries.

USE INSTRUCTIONS

Apply sufficient quantity directly on the back and / or clothes of the pet.
Ideal to be used after the usual bath.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

FOR EXTERNAL USE ONLY

CONSERVATION

Store in a cool and dry place, protected from light, at no more than 30ºC.

PRESENTATION

180 mL spray bottle

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

COMMANDER 20/20® TOPICAL SOLUTION

SOLUCIÓN TÓPICA.

ANTIPARASITARIO EXTERNO POUR-ON.

Technical Specification

SPECIES

Bovine.

Commander 20/20 Pour on, is an external antiparasitic that can be administered to cattle of any age for the control of the horn fly (*Haematobia irritans*) and lice (*Damalinia bovis, Haematopinus eurysternus* and *Linognathus vituli*). The field studies showed a maximum efficiency of up to 35 days in the control of the horn fly. In addition, it has been shown that there is an effectiveness of up to 8 weeks in the control of lice.

DOSAGE FORM

Topical solution.

THERAPEUTIC ACTION

External Antiparasitic Pour-on

COMPOSITION

Each 100 mL contains:
Permethrin (25/75)..... 20 g
Piperonyl Butoxide ..... 20 g
Excipients q.s.p..... 100 mL

MODE OF APPLICATION

With a dose gun or a syringe, apply in a dorsal longitudinal line from the base of the tail (rump) until the area of the cross. The application “against the grain” allows a better distribution of the product on the skin of the animals.

ROUTE OD ADMINISTRATION AND DOSAGE

Against fly and lice: Administer 10 mL per animal.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- The over exposure in susceptible subjects to the product, it could trigger side effects (allergies, skin rush, others)
- Use gloves when handling this product.
- In case of accidental contact, wash the affected are with soap and plentiful of water.
- In case of poisoning, call the Doctor.

WARNINGS

- Mantener fuera del alcance de los niños.
- No aplicar próximo a fuentes de calor.
- No desechar el envase en ríos, lagos u otras fuentes de agua.
- Evitar el contacto con los ojos.
- Condiciones climáticas diferentes a las presentadas en los estudios de campo pueden hacer variar la duración del efecto.

GUARD PERIOD

Meat: 0 days.

Milk: 2 days.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° and 30°C

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

100 mL, 250 mL, 1 Liter and 3 Liter flask.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 1650
- Panama: Reg. No. RF-8328-18

CONDROVET® TABLETS

COMPRIMIDO ORAL.

SUPLEMENTO NUTRICIONAL PARA PERROS.

Technical Specification

SPECIES

Dogs.

DOSAGE FORM

Tablets.

THERAPEUTIC ACTION

Nutritional supplement for dogs

COMPOSITION

Each tablet contains:

Glucosamine Sulfate.....500 mg

Chondroitin Sulfate.....400 mg

Vitamin C.....33 mg

Manganese.....5 mg

Excipients q.s.p.....1 tablet

PROPERTIES

CONDROVET® contains essential ingredients to maintain healthy joints in your dog. CONDROVET® is a natural source of the structural elements required to preserve the joint cartilage.

INDICATIONS

Do not use in animals sensitive to any of its components.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer orally as indicated in the table below:

Weight (kilogram)	Initial dose (first 6 weeks)	Maintenance dose
5 to 10	1 tablet / day	½ tablet / day
11 to 22	2 tablets / day	1 tablet / day
23 to 45	3 tablets / day	1½ tablets / day
> 45	4 tablets / day	2 tablets / day

CONTRAINDICATIONS

Do not use in animals sensitive to any of its components.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

IT IS NOT A COMPLETE FOOD.

CONSERVATION

Store in a fresh, dry place, out of direct light, at no more than 30 °C.

PRESENTATION

30 tablets.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

SAG Inscription #: RM03-010

CORTIDRAG® TOPICAL SOLUTION

SOLUCIÓN TÓPICA.

ANTIBIÓTICO – ANALGÉSICO - ANTIINFLAMATORIO ESTEROIDAL

EFICAZ CONTRA LA QUERATOCONJUNTIVITIS INFECCIOSA BOVINA

Technical Specification

SPECIES

Bovine.

Cortidrag® is recommended for the treatment of the infectious bovine karatoconjuntivitis caused by *Moraxella bovis*.

DESCRIPTION

Antibiotic – Analgesic - Steroidal anti-inflammatory

Effective against the infectious bovine keratoconjunctivitis.

DOSAGE FORM

Topical solution.

THERAPEUTIC ACTION

Antibiotic – Analgesic - Steroidal anti-inflammatory

Effective against infectious bovine keratoconjunctivitis.

COMPOSITION

Each 100 mL of topical solution contains:
Gentamicin sulphate.....0.509 g
(Equivalent to 0.30 g of basic Gentamicin)
Lidocaine Hydrochloride .....1.000 g
(Equivalent to 0.87 g of basic Lidocaine)
Dexamethasone Sodium phosphate.....0.132 g
(Equivalent to 0.10 g of Dexamethasone)
Excipients q.s.p. .....100 mL

INDICATIONS

- Do not administer in patients with hypersensitivity to any of the active ingredients.
- Do not administer in patient with renal and/or hepatic impairment.
- Do not administer to pregnant or lactating females or to breeding animals.

MODE OF APPLICATION

Clean the eye area with water and then dry. Direct the orifice of the spray valve towards the affected eye at a distance of approximately 10 cm. Apply the product by squeezing the trigger of the spray valve.

ROUTE OD ADMINISTRATION AND DOSAGE

- Topical route.
- Manage 3 sprays in each affected eye every 24 hours for 5 days.

DRUG INTERACTIONS

It is not recommended to use it concomitantly with other pharmaceutical products.

CONTRAINDICATIONS

- Do not administer in patients with hypersensitivity to any of the active ingredients.
- Do not administer in patient with renal and/or hepatic impairment.
- Do not administer to pregnant or lactating females or to breeding animals.

PRECAUTIONS

SPECIAL PRECAUTIONS FOR USE

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not handle by people hypersensitive to any active ingredient of the composition.
- Avoid inhalation of vapors.
- Do not smoke, eat or drink while handling and applying the product.
- Wash your hands with plenty of water immediately after applying the product.
- In case of contact with skin, eyes or mucous membranes, wash immediately with plenty of water.
- In case of accidental ingestion, go immediately to a medical center and show the product's packaging.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

0 days.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be discarded as domestic waste, without any special precautions. Do not dispose of containers with product residues on the ground or in watercourses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 °C, protected from light.
Once the container is opened, use within 12 weeks. Discard unused product after that time period.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Spray bottle containing 125 and 250 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No. 2202-B

CRACUL® TABLETS

COMPRIMIDO ORAL.

AYUDA A EVITAR LA INGESTA DE HECES EN PERROS.

Technical Specification

SPECIES

Dogs.

CRACUL® is a natural product for dogs, manufactured using the extract obtained from the mature, dried fruits of *Capsicum annuum* and *Capsicum frutescens L*. The extract imparts a disagreeable taste and odor to feces, discouraging its consumption by the animal.

CRACUL® is developed from an encapsulated form of highly concentrated natural extract of Capsicum, which is protected by a mixture of excipients and protective layers using a patented process, and thus, it guarantees a maximum efficacy of capsacinoids, preventing any oral or gastric irritation. Also, CRACUL® does not show any other type of side effect.

DOSAGE FORM

Tablets.

THERAPEUTIC ACTION

Help to prevent stool eating in dogs.

COMPOSITION

Each tablet contains:

Capsicum oleoresin.....40 mg

Excipients q.s.p.....1 tablet

ROUTE OD ADMINISTRATION AND DOSAGE

Administer orally, 1 tablet per each 10 kg of body weight, daily, for 2 weeks.

WARNINGS

Mantener fuera del alcance de los niños.

CONDITION OF SALE

OTC product (non-prescription)

PRESENTATION

20 tablets.

PREPARED BY

Drag Pharma Laboratory.

DALMARELIN ® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANÁLOGO SINTÉTICO DE LA HORMONA LIBERADORA DE GONADOTROPINA.

Technical Specification

SPECIES

Cows and mares.

Cows:
- It is indicated in the treatment and prevention of follicular ovarian cysts.
- Induction of ovulation at the time of insemination in cases of short, prolonged or silent heat.

Mares:
- It is indicated in the induction of ovulation.
- Improved conception rate.

DESCRIPTION

Synthetic GnRH analog

DOSAGE FORM

Injectable solution

THERAPEUTIC ACTION

Synthetic analog of gonadotropin-releasing hormone.

COMPOSITION

Each mL of solution for injection contains:
Lecirelin Acetate .....26.5 µg
(equivalent to 25 µg of Lecirelin)
Excipients csp .....1 mL

PROPERTIES

Dalmarelin® is an injectable aqueous solution containing lecirelin, a synthetic analog of the hypothalamic polopeptide hormone GnRH, which induces the release of the gonadotropin hormones LH and FSH from the frontal pituitary.
Lecirelin differs from natural hormones in that it is a nonapeptide and not a decapeptide by substituting glycine in the tenth position with a highly lipophilic ethylamine. This change in the structure increases the affinity for specific pituitary receptors, resulting in an increase in the levels of LH and FSH, as well as a prolongation of their effects for a period greater than 240 minutes, as opposed to the 90 minutes of the natural hormone.

Dalmarelin® is indicated in cases of low fertility due to immaturity or delayed maturity of the flail and ovulation due to insufficient FSH and LH secretion by the pituitary. The inoculation of gonadotropins determines physiologically uncontrollable exogenous effects, whereas the administration of **Dalmarelin®** determines the secretion of pituitary gonadotropins that can be controlled by the organism through negative feedback mechanisms.

MODE OF APPLICATION

Intramuscular administration.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer intramuscularly.

Cows

· **Treatment of follicular ovarian cysts:** 50 µg of Lecirelin per animal, equivalent to 2 mL of the product, once the existence of a cyst has been verified. If no disappearance or luteinization has occurred after 10 days, apply a second dose of 100 µg of lecirelin, equivalent to 4 mL of product.

· **Prevention of follicular ovarian cysts:** In cows with a history of previous lactations of follicular cysts, apply 50 µg of Lecirelin per animal, equivalent to 2 mL of the product, on day 14 post partum.

· **Induction of ovulation at the time of insemination in cases of short, prolonged or silent cycles:** 50 µg of Lecirelin per animal, equivalent to 2 mL of the product at the time of insemination.

Mares

· **Induction of ovulation and improvement of the conception rate:** 4 mL of product per animal. Administration should be carried out when, after gynecological inspection, a follicle of at least 40 mm in diameter is diagnosed.
The treatment should be repeated if after a period of 24-36 hours there is no ovulation.

WARNINGS

- Su uso no está recomendado durante la gestación.
- En caso de contacto con la piel del operador, se recomienda lavar inmediatamente con abundante agua y jabón.
- No manipular por mujeres embarazadas.

GUARD PERIOD

Meat: Zero days.

Milk: Zero hours.

CONSERVATION

Store between 2 and 25 ° C in a cool, dry place protected from light.

Once opened, use within 28 days. Discard the unused product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Case with 1 or 5 bottles of 10 mL, 1 or 10 bottles of 4 mL or 1 bottle of 20 mL of product

PREPARED BY

Fatro S.p.A.- Italy.

RECORDS

Reg SAG No 2077-B

D-COMPLEX® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

VITAMÍNICO RECONSTITUYENTE.

Technical Specification

SPECIES

Horses.

Vitamin B complex, general restorative. Appetite stimulant, neurobiotic, liver protector and lipotropic. Recommended in pictures of vitamin deficiency of the B complex; as support therapy in animals convalescing from parasitic, infectious or surgical diseases; in the treatment of neuritis, neuralgia and neuropathies in general. It is also recommended in states of higher nutritional requirement, such as: pregnancy, lactation, growth and sports training.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Vitaminic strengthening.

COMPOSITION

Each mL contains:
Thiamine Hydrochloride .....150 mg
Riboflavin 5-Sodium phosphate.....2 mg
Pyridoxine Hydrochloride.....10 mg
Nicotinamide.....150 mg
D-panthenol.....10 mg
Choline Chloride.....20 mg
Inositol.....20 mg
Excipients q.s.p.....1 mL

INDICATIONS

Do not use in animals with hypersensitivity to thiamine administered parenterally, because of the risk of anaphylactic shock.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration route: Intramuscular o intravenous slowly.

Product dose:

5 mL per animal, once a week for up to 4 weeks.

CONTRAINDICATIONS

Do not use in animals with hypersensitivity to thiamine administered parenterally, because of the risk of anaphylactic shock.

PRECAUTIONS

- An anaphylactic reaction may occur in individuals who are hypersensitive to parenteral Thiamine; possible risk of anaphylactic shock.

- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR USE

- An anaphylactic reaction may occur in individuals who are hypersensitive to parenteral Thiamine; possible risk of anaphylactic shock.

- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not handle by people who are hypersensitive to Thiamine or any of its components.
- In case of contact with skin, eyes or mucous membranes, wash immediately with plenty of water.
- In case of accidental injection, go immediately to a medical center and show the product label.

GUARD PERIOD

Zero (0) days.

OBSERVATIONS

Special precautions for disposing of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° y 30°C, protected from sunlight.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

100 mL vial

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG No. 1588

DEPODRAG® EQUINE - INJECTABLE SUSPENSION

SUSPENSIÓN INYECTABLE.

ANTIINFLAMATORIO ESTEROIDAL.

Technical Specification

SPECIES

Horses

Anti-inflammatory of long lasting effect for allergic, dermatologic disorders and arthritis.

DOSAGE FORM

Injectable suspension.

THERAPEUTIC ACTION

Steroidal anti-inflammatory.

COMPOSITION

Each 1 mL of suspension contains:
Triamcinolone Acetonide..... 6 mg
Excipients q.s.p.....1 mL

INDICATIONS

- Do not use in viral infectious processes and generalized fungal infections.
- Do not use in musculoskeletal disorders where immobility is required.
- Do not use in animals with tuberculosis, chronic nephritis or Cushing's syndrome, except for emergency therapy.
- Do not use in case of bone metaplasia and osteoporosis.
- Do not use in treatment of laminitis.
- Do not use in patients with diabetes mellitus, kidney or heart failure.
- Do not use in patients with hypersensitivity to the active substance.
- Do not use in animals with gastrointestinal or corneal ulcers.
- Do not use in pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration route: Intramuscular, subcutaneous, intra-articular and intrasynovial.

Recommended dose:
Horses:

- Intramuscular or subcutaneous route: 2 to 5 mL in a single dose.
- Intra-articular or intrasynovial route: 1 to 5 mL in a single dose. Repeat if necessary under Veterinary Medical recommendation.

It is not recommended to repeat the treatment beyond 3 consecutive times.

DRUG INTERACTIONS

- Amphotericin B or caliuretic diuretics (Furosemide, Thiazides) can cause hypokalemia when administered concomitantly with glucocorticoids. When these drugs are used simultaneously with digitalis glycosides, it may increase the possibility of toxicity if hypokalemia is generated. Diligent monitoring of potassium and digitalis levels is recommended.
- Glucocorticoids can lower blood levels of salicylates.
- Insulin requirements may increase in patients receiving glucocorticoids.
- Phenytoin, Phenobarbital, Rifampicin can increase glucocorticoid metabolism.
- Concomitant administration of glucocorticoids and cyclosporine can increase the blood levels of each, with mutual inhibition of liver metabolism. The clinical importance of this interaction is uncertain. Glucocorticoids can also inhibit hepatic metabolism of Cyclophosphamide. Dosage adjustments may be required.
- Mitotane can alter steroid metabolism; higher doses than usual may be necessary to treat mitotane-induced adrenal insufficiency.
- Patients treated with corticosteroids at immunosuppressive doses should not receive live attenuated live virus vaccines because viral replication may be enhanced. A decreased immune response may occur after administration of a vaccine, toxoid, or bacterin, in patients receiving glucocorticoids.
- Administration of ulcerogenic drugs (eg, nonsteroidal anti-inflammatory drugs) with glucocorticoids may increase the risk of gastrointestinal ulceration.
- The effects of Hydrocortisone, and possibly other glucocorticoids, can be potentiated by concomitant administration with estrogens.
- In patients with myasthenia gravis, concomitant administration of glucocorticoids and anticholinesterases (eg, Pyridostigmine, Neostigmine, etc.) can induce pronounced muscle weakness. If possible, discontinue anticholinesterase medication for at least 24 hours before glucocorticoid administration.

CONTRAINDICATIONS

- Do not use in viral infectious processes and generalized fungal infections.
- Do not use in musculoskeletal disorders where immobility is required.
- Do not use in animals with tuberculosis, chronic nephritis or Cushing's syndrome, except for emergency therapy.
- Do not use in case of bone metaplasia and osteoporosis.
- Do not use in treatment of laminitis.
- Do not use in patients with diabetes mellitus, kidney or heart failure.
- Do not use in patients with hypersensitivity to the active substance.
- Do not use in animals with gastrointestinal or corneal ulcers.
- Do not use in pregnant or lactating females.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- The product is irritating in the case of contact with the eyes. It can be dangerous in the case of accidental ingestion and in the case of contact with the skin.
- Pregnant women should not handle the product.

ADVERSE EFFECTS

Unwanted effects and adverse reactions:
Prolonged use of Depodrag® can cause suppressive effects on the hypothalamic-pituitary-adrenal axis leading to adrenal atrophy (adrenal insufficiency). It can also cause bone resorption or inhibition of growth and bone repair, inhibition of collagen synthesis, decreased growth rate, delayed healing, diarrhea, gastrointestinal irritation, gastrointestinal ulceration, hematopoietic changes, retention of sodium and fluid and flares of latent infections.
The most frequent side effects are polyuria, polydipsia, polyphagia, lethargy, weakness and bilateral alopecia. Less frequent are weight loss, anorexia, and diarrhea.

GUARD PERIOD

Do not use in horses intended for human consumption.

OBSERVATIONS

Use during pregnancy, lactation and in breeding animals:

- Do not use in pregnant and lactating females.
- Glucocorticoids are probably necessary for normal fetal development.
- They may be required for adequate surfactant production and development of myelin, retina, pancreas, and breasts.
- Excessive doses early in pregnancy can lead to teratogenic effects. In horses, the administration of exogenous steroids can induce labor when used in late pregnancy. It is recommended not to use high doses in pregnant animals.
- Glucocorticoids not bound to plasma proteins enter milk. High doses or prolonged administration in mothers can potentially inhibit the growth of newborns.

Special precautions for the disposal of unused product or waste material:
Discard the remains of unused product in its original container. Dispose of this product with caution with household waste.
Uruguay: Dispose of the product container at the nearest collection center

CONSERVATION

Store at a temperature between 2 ° and 30° C, protected from light. Once opened, use within 3 months. Discard the unused product after that period of time.

CONDITION OF SALE

To be supply only with veterinary prescription.

PRESENTATION

Ampule containing 20 mL in a box with 3 ampules containing 5 mL each.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG Nº 0529
- Bolivia: Reg. SENASAG PUV-F-N° 005515/13
- Uruguay: Reg. MGAP N° A-4493
- Rep. Dominicana: Reg. Nº 5607
- Perú: Reg. SENASA F.06.42.I.0240

DEPODRAG® PET - INJECTABLE SUSPENSION

SUSPENSIÓN INYECTABLE.

ANTIINFLAMATORIO ESTEROIDAL.

Technical Specification

SPECIES

Dogs and cats.

Anti-inflammatory of long lasting effect for allergic and dermatologic disorders and arthritis.

DOSAGE FORM

Injectable suspension.

THERAPEUTIC ACTION

Steroidal anti-inflammatory.

COMPOSITION

Each 1 mL of suspension contains:
Triamcinolone Acetonide..... 6 mg
Excipients q.s.p.....1 mL

INDICATIONS

- Do not use in viral infectious processes and generalized fungal infections.
- Do not use in musculoskeletal disorders where immobility is required.
- Do not use in animals with tuberculosis, chronic nephritis or Cushing's syndrome, except for emergency therapy.
- Do not use in case of bone metaplasia and osteoporosis.
- Do not use in patients with diabetes mellitus, kidney or heart failure.
- Do not use in patients with hypersensitivity to the active substance.
- Do not use in animals with gastrointestinal or corneal ulcer.
- Do not use in pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration route: Intramuscular, subcutaneous, intra-articular e intrasynovial.

Dose of the active principle:
Dogs and cats:
- In allergy symptoms: 0.2 mg / Kg, in a single dose. In severe cases, it can be administered up to 1 mg / Kg, in a single dose.
- In intra-articular and intrasynovial treatment: 1 to 3 mg / Kg, in a single dose, repeated if necessary after 3 to 4 days.

Product dosage:
Dogs and cats:
- In allergy : 0.2 mL every 6 Kg of weight, in a single dose. In severe cases, up to 1 mL can be administered every 6 kg of weight, in a single dose.
- In intra-articular and intrasynovial treatment: 0.2 to 0.5 mL, in a single dose, repeated if necessary after 3 to 4 days.
It is not advisable to repeat the treatment beyond three consecutive times.

DRUG INTERACTIONS

- Amphotericin B or caliuretic diuretics (Furosemide, Thiazides) can cause hypokalaemia when administered concomitantly with glucocorticoids. When these drugs are used simultaneously with digitalis glycosides, the possibility of toxicity may be increased if hypokalemia develops. Diligent monitoring of potassium and digitalis levels is recommended.
- Glucocorticoids can lower blood levels of salicylates.
- Insulin requirements may increase in patients receiving glucocorticoids.
- Phenytoin, Phenobarbital, Rifampicin can increase glucocorticoid metabolism.
- The concomitant administration of glucocorticoids and cyclosporine can increase the blood levels of each, with mutual inhibition of hepatic metabolism. The clinical significance of this interaction is uncertain. Glucocorticoids can also inhibit the hepatic metabolism of cyclophosphamide. Dosage adjustments may be required.
- Mitotane can alter steroid metabolism; Higher than usual doses may be necessary to treat mitotane-induced adrenal insufficiency.
- Patients treated with corticosteroids at immunosuppressive doses should not receive live attenuated live virus vaccines because viral replication may be enhanced. A lowered immune response can occur after administration of a vaccine, toxoid, or bacterin, in patients receiving glucocorticoids.
- Administration of ulcerogenic drugs (eg, non-steroidal anti-inflammatory drugs) with glucocorticoids may increase the risk of gastrointestinal ulceration.
- The effects of hydrocortisone, and possibly other glucocorticoids, can be potentiated by concomitant administration with estrogens.
- In patients with Myasthenia gravis, concomitant administration of glucocorticoids and anticholinesterases (eg, Pyridostigmine, Neostigmine, etc.) can induce pronounced muscle weakness. If possible, discontinue anticholinesterase medication for at least 24 hours before glucocorticoid administration.

CONTRAINDICATIONS

- Do not use in viral infectious processes and generalized fungal infections.
- Do not use in musculoskeletal disorders where immobility is required.
- Do not use in animals with tuberculosis, chronic nephritis or Cushing's syndrome, except for emergency therapy.
- Do not use in case of bone metaplasia and osteoporosis.
- Do not use in patients with diabetes mellitus, kidney or heart failure.
- Do not use in patients with hypersensitivity to the active substance.
- Do not use in animals with gastrointestinal or corneal ulcer.
- Do not use in pregnant or lactating females.

PRECAUTIONS

Special warnings and precautions for use:

- In bacterial infections, the use must be associated with antibacterials.
- Corticosteroids can precipitate labor during the final stages of pregnancy.
- Shake before using.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR USE

Special warnings and precautions for use:

- In bacterial infections, the use must be associated with antibacterials.
- Corticosteroids can precipitate labor during the final stages of pregnancy.
- Shake before using.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- The product is irritating in the case of contact with the eyes. It can be dangerous in the case of accidental ingestion and in the case of contact with the skin.
- Pregnant women should not handle the product.

ADVERSE EFFECTS

Unwanted effects and adverse reactions:

Long-term use of Depodrag® may cause suppressive effects on the hypothalamic-pituitary-adrenal axis leading to adrenal atrophy (adrenal insufficiency). It can also cause bone resorption or inhibition of bone growth and repair, inhibition of collagen synthesis, decreased growth rate, delayed healing, diarrhea, gastrointestinal irritation, gastrointestinal ulceration, hematopoietic changes, sodium and sodium retention. fluid and flare-up of latent infections.
The most common side effects are polyuria, polydipsia, polyphagia, lethargy, weakness, and bilateral alopecia. Less common are weight loss, anorexia, and diarrhea.

OBSERVATIONS

Do not use in pregnant and lactating females.

- Glucocorticoids are probably necessary for normal fetal development. They may be required for proper surfactant production and development of myelin, retina, pancreas, and breasts.
- Excessive doses early in gestation can lead to teratogenic effects. The administration of exogenous steroids can induce labor when used in the final stages of pregnancy. It is recommended not to use high doses in pregnant animals.
- Glucocorticoids not bound to plasma proteins enter the milk. High doses or prolonged administration to mothers can potentially inhibit the growth of newborns.

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Dispose of the waste of this product with care together with household waste.
Uruguay: Dispose of the product container at the nearest collection center.

CONSERVATION

Store at a temperature between 2 ° and 30 ° C, protected from light. Once opened use within 3 months. Discard the unused product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

5 mL and 20 mL vial

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG Nº 0529
- Uruguay: Reg. N° MGAP A-4493
- Rep. Dominicana: Reg. Nº 5607
- Panamá: Reg. Nº RF-4183-19
- Bolivia: Reg. SENASAG CR-PUV Nº 005515/13
- Perú: Reg. SENASA F.06.42.I.0240

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

Imported in Uruguay:

VIVAFIL S.A.
Rio Negro 1107 Montevideo - Uruguay, TEL 29001112
grupotecnovet@gmail.com
Technical Director: DMTV Diego Cuadrado.

Imported and distributed in Peru by:

Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

DERMISOLONA® ORAL SUSPENSION

SUSPENSIÓN ORAL.

ANTIINFLAMATORIO – ANTIALÉRGICO ESTEROIDAL.

Technical Specification

SPECIES

Dogs and cats.

It is indicated as an aid in the treatment of inflammatory and allergic disorders in cats and dogs.

In dogs it is also indicated as an immunosuppressant in autoimmune diseases, as a therapy for chronic inflammatory bowel disease and in the treatment of hypoadrenocorticism.

DOSAGE FORM

Oral suspension

THERAPEUTIC ACTION

Anti-inflammatory - Steroidal antiallergic

COMPOSITION

Each mL of oral suspension contains:
Prednisolone Acetate ..... 4.5 mg
(Equivalent to 4 mg of Prednisolone base)
Excipients q.s. ..... 1 mL

INDICATIONS

- Do not use in animals with Cushing's syndrome.
- Do not use in patients with systemic fungal infections or viral infections.
- Do not use in animals with peptic or corneal ulcers.
- Do not administer to pregnant females.
- Do not use in animals hypersensitive to any of the components of this product.
- Do not administer in conjunction with vaccines.

USE INSTRUCTIONS

Shake the suspension before use, then withdraw the product dose with the dosing syringe or dropper and administer directly into the oral cavity.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration

	DOSE OF THE ACTIVE SUBSTANCE	PRODUCT DOSE (1 mL = 26 drops of product)
ANTI-INFLAMATORY AND ANTI-ALLERGIC	DOGS 0,5 to 1 mg/Kg of weight, every 12 a 24 hours, for 3 to 5 days. CATS 1 to 2 mg/Kg of weight, every 12 to 24 hours, for 3 to 5 days.	1,5 to 2,5 mL for each 10 Kg of weight, or 4 to 7 gotas for each Kg of weight,, for each 12 to 24 hours, for 3 to 5 days. 1,5 to 2,5 mL for every 5 Kg of weight or 7 to 13 drops for each Kg of weight, every 12 to 24 hours, for 3 to 5 días.
IMMUNOSUPPRESSOR	In prolonged treatments, it is recommended to reduce to the lowest effective dose and administer on alternate days in the morning for dogs, and in the afternoon in cats (to respect the circadian rhythm). DOGS 1 to 4 mg / Kg / every 24 hours for 2 to 14 days. Then, 0.5 to 2 mg / Kg / every 24 hours, for 14 days. Then reduce the dose at regular intervals (2 to 6 weeks), until reaching the maintenance dose of 0.5 to 1 mg / Kg / every other day. (*)	2.5 to 10 mL for every 10 Kg of weight per day, or 7 to 26 drops for each Kg of weight every 24 hours, for 2 to 14 days. Then, 1.3 to 5 mL for every 10 Kg of weight per day, or 4 to 13 drops for each Kg of weight every 24 hours, for 2 to 14 days. Then reduce the dose at regular intervals (2 to 6 weeks), until reaching the maintenance dose of 1.5 to 2.5 mL for every 10 Kg of weight, or 4 to 7 drops for each Kg of weight, every other day . (*) (**)
CHRONIC INFLAMMATORY INTESTINAL DISEASE	DOGS 1 to 4 mg / Kg of weight every 12 to 24 hours, for 2 to 4 weeks, then decrease the dose slowly at intervals of 10 days to 2 weeks depending on the clinical response, until reaching a maintenance dose of 0.5 to 1 mg / Kg every other day (*)	2.5 to 10 mL for every 10 Kg of weight per day, or 7 to 26 drops for each Kg of weight, every 12 to 24 hours, for 2 to 4 weeks, then decrease the dose slowly at intervals of 10 days to 2 weeks depending on the clinical response, until reaching the maintenance dose of 1.5 to 2.5 mL for every 10 Kg of weight, or 4 to 7 drops for each Kg of weight, on alternate days. (*) (**)
HYPOADRENOCORTICISM	DOGS 0.2 to 0.4 mg / Kg of weight, every 24 hours, permanently.	0.5 to 1 mL per 10 Kg of weight or 2 to 3 drops per Kg of weight every 24 hours, permanently.

(*) The duration of therapy varies according to the intensity of the symptoms and the remission of symptoms, and can last for several weeks or months. In these cases, it is recommended to use the minimum

effective dose, every other day.

(**) 1 mL of DERMISOLONA® Oral suspension is equivalent to 26 drops of product.

DRUG INTERACTIONS

It is recommended to avoid the concomitant administration of Prednisolone with:

- Amphotericin B or caliuretic diuretics (furosemide, thiazides), as hypokalemia may occur. Digitalis may increase the possibility of toxicity, if hypokalemia is generated.
- Phenytoin, phenobarbital and rifampicin, because they can increase the metabolism of corticosteroids.
- Cyclosporine, because the blood levels of both drugs can be increased with mutual inhibition of liver metabolism.
- With ulcerogenic drugs (for example, non-steroidal anti-inflammatory drugs) since the risk of gastrointestinal ulceration can be increased.
- Cyclophosphamide, as corticosteroids can inhibit hepatic chemotherapy metabolism.
- Patients treated with corticosteroids at immunosuppressive doses generally should not receive live attenuated virus vaccines, because viral replication may be enhanced.
- Insulin requirements may be increased in patients receiving glucocorticoids.

CONTRAINDICATIONS

- Do not use in animals with Cushing's syndrome.
- Do not use in patients with systemic fungal infections or viral infections.
- Do not use in animals with peptic or corneal ulcers.
- Do not administer to pregnant females.
- Do not use in animals hypersensitive to any of the components of this product.
- Do not administer in conjunction with vaccines.

PRECAUTIONS

- In case of prolonged treatments with Prednisolone, a higher protein intake must be provided to keep the animal in a positive nitrogen balance.
- No delayed effect on wound healing has been described, however, such a possibility should be considered when used in surgery.
- Prolonged glucocorticoid therapy can suppress adrenocortical activity, so discontinuation should be done gradually to ensure gradual return of ACTH and endogenous corticosteroid functions.
- Administer with caution in animals that have diabetes, osteoporosis, or are recovering from a bone fracture, predisposition to thrombophlebitis, hypertension, congestive heart failure, kidney failure, or active tuberculosis.
- The anti-inflammatory effects can mask signs of infection.
- Administer with caution in growing animals as it may cause delay.
- Administer with caution in animals with liver failure.

SPECIAL PRECAUTIONS FOR USE

- In case of prolonged treatments with Prednisolone, a higher protein intake must be provided to keep the animal in a positive nitrogen balance.
- No delayed effect on wound healing has been described, however, such a possibility should be considered when used in surgery.
- Prolonged glucocorticoid therapy can suppress adrenocortical activity, so discontinuation should be done gradually to ensure gradual return of ACTH and endogenous corticosteroid functions.
- Administer with caution in animals that have diabetes, osteoporosis, or are recovering from a bone fracture, predisposition to thrombophlebitis, hypertension, congestive heart failure, kidney failure, or active tuberculosis.
- The anti-inflammatory effects can mask signs of infection.
- Administer with caution in growing animals as it may cause delay.
- Administer with caution in animals with liver failure.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Wash hands after administering the product.
- In case of skin contact, it is recommended to wash your hands with soap and plenty of water. If irritation develops and persists, see a doctor.
- In the case of contact with the eyes, it is recommended to wash with plenty of water for 15 minutes. If irritation develops and persists, see a doctor.
- In case of ingestion, do not induce vomiting. Get medical help.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

- Administered in the short term, Prednisolone is unlikely to cause harmful effects, even in massive doses. Adverse effects are generally associated with chronic therapies, especially at high doses or if an alternate day regimen is not followed.
- The most frequent adverse effects are polyuria, polydipsia and polyphagia; and they are preferably associated with anti-inflammatory doses.
- With regard to immunosuppressive doses and extensive treatments, adverse reactions are more likely to appear and are potentially more pronounced. These effects, rarely observed, are manifested as symptoms of hyperadrenocorticism (Cushing's syndrome): opaque and dry fur, weight gain, panting, vomiting, diarrhea, hepatomegaly with the consequent alteration of the concentration of liver enzymes in the serum, pancreatitis, gastrointestinal ulceration (particularly when used with NSAIDs), lipidemias, alteration of insulin requirements, activation or intensification of Diabetes Mellitus, muscle wasting and changes in behavior (depression, lethargy, vices).
- In prolonged treatments, an increase in the incidence of osteoporosis is described, especially in elderly dogs.

OBSERVATIONS

Overdose:
Given in the short term, glucocorticoids are unlikely to cause harmful effects, even in massive doses. Adverse effects are generally associated with chronic therapies, especially at high doses or if an alternate day regimen is not followed.

Special precautions for the disposal of unused product or waste material:
Discard the remains of unused product in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of natural water. Dispose of this product with caution with household waste.

CONSERVATION

Store in a cool and dry place, at a temperature between 2 and 30º C.
Once the bottle is opened, use within 30 days.

CONDITION OF SALE

Sale with Veterinary Medical prescription only

PRESENTATION

Bottle with 30 mL

PREPARED BY

Laboratorio Drag Pharma.

RECORDS

Reg. SAG N° 2281

DERMISOLONA® ORAL TABLETS

COMPRIMIDO ORAL.

ANTIINFLAMATORIO - ANTIALÉRGICO ESTEROIDAL.

Technical Specification

SPECIES

Dogs and cats.

It is indicated as an aid in the treatment of non-infectious inflammatory disorders, non-septic arthropathies and in the treatment of allergic dermatitis; in chronic inflammatory bowel disease and as an immunosuppressant in tumor states.

DOSAGE FORM

Oral tablet.

THERAPEUTIC ACTION

Anti-inflammatory – Steroidal anti-allergic.

COMPOSITION

Each Tablet contains:
Prednisolone base.....20 mg
Excipients q.s.p.....1 tablet

INDICATIONS

- Do not use in patients with systemic fungal infections or viral infections.
- Systemic therapy with Prednisolone is contraindicated in animals with peptic ulcer, corneal ulcer, and Cushing's syndrome.
- Do not administer to pregnant animals.
- Do not use in animals hypersensitive to some of the components of this product.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration.

Active ingredient dose:

- **For anti-inflammatory and anti-allergic treatment:** 0.5 to 1 mg/Kg in dogs and 1 to 2 mg/Kg in cats, every 12 or 24 hours for 3 to 5 days. In the event of prolonged treatment, it is recommended to reduce to the lowest effective dose and give in the regime of alternated days in the morning for dogs and in the afternoon in cats (to respect the circadian rhythm).
- **As immunosuppressant:** 2.2 to 3.3 mg/Kg of weight for two consecutive days and then 2-4 mg/Kg every 48 hours. Evaluate the duration of the treatment according to the intensity and remission of the symptoms.
- **Chronic inflammatory bowel disease:** 1-2 mg/Kg of weight once a day for 2 to 4 weeks.

Product dose:

- **As anti-inflammatory and anti-allergic:**
  Dogs: ¼ - ½ tablet/10 Kg of weight every 12 or 24 hours for 3 to 5 days.
  Cats: ¼-½ Tablet /5 Kg of weight every 12 or 24 hours for 3 to 5 days.
- **As immunosuppressant:** 1 - 1 ½ tablet/10 Kg of weight for two consecutive days and then 1 - 2 tablets/10 Kg of weight, every 48 hours. Evaluate the duration of the treatment according to intensity and remission of the symptoms.
- **Chronic inflammatory bowel disease:** ½ - 1 tablet/10 Kg of weight once a day for 2 to 4 weeks.

DRUG INTERACTIONS

It is recommended to avoid the concomitant administration of Prednisolone with:

- Amphotericin B or caliuretic diuretics (Furosemide, Thiazides), as hypokalemia may occur. Digitalis may increase the possibility of toxicity, if hypokalemia is generated.
- Phenytoin, Phenobarbital and Rifampicin, because they can increase the metabolism of corticosteroids.
- Cyclosporine, because the blood levels of both drugs can be increased with mutual inhibition of liver metabolism.
- With ulcerogenic drugs (for example, non-steroidal anti-inflammatory drugs) since the risk of gastrointestinal ulceration can be increased.
- Cyclophosphamide, as corticosteroids can inhibit hepatic chemotherapy metabolism.
- Patients treated with corticosteroids at immunosuppressive doses generally should not receive live attenuated virus vaccines, because viral replication may be enhanced.
- Insulin requirements may be increased in patients receiving glucocorticoids.

CONTRAINDICATIONS

- Do not use in patients with systemic fungal infections or viral infections.
- Systemic therapy with Prednisolone is contraindicated in animals with peptic ulcer, corneal ulcer, and Cushing's syndrome.
- Do not administer to pregnant animals.
- Do not use in animals hypersensitive to some of the components of this product.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Wash hands after administering the product

WARNINGS

Advertencias y precauciones especiales de uso:

- En caso de tratamientos prolongados con Prednisolona se debe proporcionar una mayor ingesta de proteínas para mantener al animal en un balance de Nitrógeno positivo.
- No ha sido descrito un efecto de retraso en la cicatrización de heridas, sin embargo, tal posibilidad debe ser considerada al ser utilizado en cirugía.
- La terapia prolongada con glucocorticoides puede suprimir la actividad adrenocortical, de manera que la suspensión de la misma debe realizarse en forma graduada para asegurar el retorno paulatino de las funciones de ACTH y corticoides endógenos.
- Administrar con precaución en animales que presenten diabetes, osteoporosis, o que estén recuperándose de una fractura ósea, predisposición a la tromboflebitis, hipertensión, insuficiencia cardiaca congestiva, insuficiencia renal o tuberculosis activa.
- Los efectos antinflamatorios pueden enmascarar signos de infección.
- Administrar con precaución en animales en crecimiento ya que puede causar retardo.
- Administrar con precaución en animales con insuficiencia hepática.
- La Prednisolona pasa a la leche y podría inducir efectos negativos en los lactantes, pero solo si la madre recibe dosis elevadas durante períodos prolongados.
- Mantener alejado del alcance de los niños.

SIDE EFFECTS

- Therapeutic use of Prednisolone is unlikely to cause metabolic-like effects associated with glucocorticoids. An effect of delayed wound healing has not been described, however, such a possibility should be considered when used in surgery.
- Long-term treatment describes an increase in the incidence of osteoporosis, especially in older dogs. Its use is not recommended during the recovery phase of a bone fracture.
- The most frequent adverse effects are polyuria, polydipsia and polyphagia; and they are preferably associated with anti-inflammatory doses.
- Regarding immunosuppressive doses and extensive treatments, adverse reactions are more likely to occur and are potentially more pronounced. Such effects, which are rarely observed, are manifested as symptoms of hyperadrenocorticism (Cushing's syndrome): opaque and dry fur, weight gain, panting, vomiting, diarrhea, hepatomegaly with the consequent alteration of the concentration of liver enzymes in the serum, pancreatitis, gastrointestinal ulceration (particularly when used with NSAIDs), lipidemias, altered insulin requirements, activation or intensification of diabetes mellitus, muscle wasting, and behavioral changes (depression, lethargy, vices).

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:

- Discard the remains of unused product in its original container.
- Do not throw the empty container or with product remains, in rivers, lakes or streams of natural water.
- Dispose of this product with caution together with household waste.
- Contact the manufacturing company or companies specialized in waste disposal, to receive recommendations on the disposal of expired or unused products.

CONSERVATION

Store at a temperature between 15 and 30°C, protected from light.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Box containing 10 tables.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG N° 1745
Bolivia: Reg. SENASAG PUV-F-N° 005507/13

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

DF-7® DRENCH - ORAL SUSPENSION

SUSPENSIÓN ORAL.

ANTIPARASITARIO INTERNO DE AMPLIO ESPECTRO.

Technical Specification

SPECIES

Horses.

Internal anti-parasitic of wide spectrum in single dose. Recommended for the treatment of parasitic nematodes gastrointestinal and migratory in adult and larval state. Also is effective against *Gasterophillus spp.* and *Anoplocephala perfoliata* in horses.

DOSAGE FORM

Oral suspension.

THERAPEUTIC ACTION

Internal anti-parasitic of wide spectrum.

COMPOSITION

Each 100 mL of suspension contains:
Ivermectin.....0.200 g
Pyrantel Pamoate.....28.53 g
(Equivalent to 9.9 g of Pyrantel base)
Excipients: q.s.....100 mL

INDICATIONS

- Do not use in horses less than 4 weeks of age.
- Do not administer to horses with known hypersensitivity to any of the active ingredients.
- Do not use in pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration.

Active ingredients dose:

- Pyrantel: 9.9 mg/Kg of weight.
- Ivermectin: 0.2 mg/Kg of weight.

Product dose:

- 1 mL of suspension for every 10 Kg of weight.

CONTRAINDICATIONS

- Do not use in horses less than 4 weeks of age.
- Do not administer to horses with known hypersensitivity to any of the active ingredients.
- Do not use in pregnant or lactating females.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Do not eat, drink or smoke while handling the product.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

Unwanted effects and adverse reactions:
The components of the df-7 drench, Oral Suspension, have a wide margin of safety. However, in case of high infestations, the treatment can generate transient colic and loose stools as a result of the parasitic destruction.

GUARD PERIOD

Do not administer to animals intended for human consumption.

OBSERVATIONS

SHAKE WELL BEFORE USE

Because it is a suspension, it should be shaken before and during administration, to maintain an optimal homogenization. Pleasant apple flavor.

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store in a cool and dry place, at room temperature between 15 and 30 ° C, protected from light.
Once opened, use the product within 12 weeks.
Discard the unused product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Bottle with 1 Liter.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 1521-B
- Dominican Republic: Reg. No. 5600

DF-7® ORAL PASTE

PASTA ORAL.

ANTIPARASITARIO INTERNO DE AMPLIO ESPECTRO

Technical Specification

SPECIES

Horses.

Internal anti-parasitic of wide spectrum in single dose. Recommended for the treatment of parasitic nematodes gastrointestinal and migratory in adult or larval state. Also is effective against *Gasterophilus spp*. and *Anoplocephala perfoliata* in horses.

DOSAGE FORM

Oral paste.

THERAPEUTIC ACTION

Internal anti-parasitic of wide spectrum.

COMPOSITION

Each syringe with 40 g of paste contains:
Ivermectin.....0.12 g
Pyrantel pamoate.....17.11 g
(Equivalent to 5.94 g of Pyrantel)
Excipients q.s.p.....40.00 g

INDICATIONS

- Do not use in horses under 4 weeks of age.
- Do not use during pregnancy or lactation, or in breeding animals.

ROUTE OD ADMINISTRATION AND DOSAGE

Dosage of active ingredients:
9.9 mg of Pyrantel and 0.2 mg of Ivermectin / Kg of body weight.

Product dosage:
Administer orally, one mark on the graduated syringe (10 grams) for every 150 kilos of body weight.
In case of high infestation, repeat the treatment according to veterinary medical advice.
40 grams of D.F. 7® pasta is enough to treat 600 kilos of weight.

DRUG INTERACTIONS

Do not use simultaneously with Morantel or Levamisole.

CONTRAINDICATIONS

- Do not use in horses under 4 weeks of age.
- Do not use during pregnancy or lactation, or in breeding animals.

PRECAUTIONS

Warnings and precautions for use:

- Keep out of the reach of children.
- Do not eat, drink or smoke while handling the product.

SPECIAL PRECAUTIONS FOR USE

Warnings and precautions for use:

- Keep out of the reach of children.
- Do not eat, drink or smoke while handling the product.

ADVERSE EFFECTS

Unwanted effects and adverse reactions:

The components of D.F. 7® pasta have a wide margin of safety. However, in case of high infestations, the treatment can generate transient colic and loose stools as a result of the parasitic destruction.

GUARD PERIOD

Do not administer to animals intended for human consumption.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of water. Dispose of the waste of this product with care together with household waste. Contact the manufacturing company or companies specialized in waste disposal, to receive recommendations on the disposal of expired or unused products.

CONSERVATION

Store at room temperature between 15 and 30 ° C.
Use immediately once opened and discard the excess product.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Box with one syringe containing 40 grams

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- **Chile:** Reg. SAG Nº 0487-B
- **Rep. Dominicana:** Reg. Nº 5601
- **Perú:** Registro SENASA F.08.43.I.1069

COUNTRIES WHERE IT IS MARKETED

Imported and Distributed in Peru by Representaciones Durand SAC.
Av. Manuel Olguín N° 501 Oficina N° 604 Santiago de Surco Lima.

DIARREPAS® ORAL SUSPENSION

SUSPENSIÓN ORAL.

ANTIDIARREICO Y QUIMIOTERÁPICO.

Technical Specification

SPECIES

Dogs and cats.

It is recommended for use in intestinal infections caused by sensitive to non-absorbable sulfonamides organisms. It is recommended for the treatment of enteric infections caused by gram positive and gram negative organisms. It has therapeutic use in acute and chronic diarrhea cases in dogs and cats, since it has protective mucosal chemotherapeutic action that avoids injury in the intestinal epithelium caused by irritating substances. By the presence of intestinal adsorbents, also prevent the absorption of toxins, gases or other toxic substances of organic or inorganic origins that can complicate a diarrheal case.

DOSAGE FORM

Oral suspension.

THERAPEUTIC ACTION

Antidiarrheal and chemotherapy.

COMPOSITION

Each 100 mL contains:
Phthalylsulfathiazole .....10 g
Sulfaguanidine.....10 g
Colloidal kaolin.....20 g
Citrus pectin .....1 g
Excipients q.s.p.....100 mL

INDICATIONS

- Do not use simultaneously or associated with other chemotherapeutic agents, or other substances with similar action.
- Do not administer to animals showing resistance to the sulfonamides treatment.
- It should not be used in animals presenting a history of hypersensitivity to some component of the product.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route.

- **Dosage of the active ingredients:** Phthalsulfatiazole 100 mg / Kg; Sulfaguanidine 100 mg / Kg; Colloidal Kaolin 200 mg / Kg, Citric Pectin 10 mg / Kg, every 12 hours for 4 days.
- **Product dose:** 10 mL for every 10 kilos of weight, every 12 hours for 4 days.

CONTRAINDICATIONS

- Do not use simultaneously or associated with other chemotherapeutic agents, or other substances with similar action.
- Do not administer to animals showing resistance to the sulfonamides treatment.
- It should not be used in animals presenting a history of hypersensitivity to some component of the product.

WARNINGS

- Mantener fuera del alcance de los niños.
- Administrar con precaución en pacientes con insuficiencia renal o hepática.
- Usar con precaución en estados de preñez avanzada.

OBSERVATIONS

Shake before using.

Special precautions for the disposal of unused product or waste material:
Discard the remains of unused product in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of natural water. Dispose of this product with caution with household waste.

CONSERVATION

Store at a temperature between 15 and 30 ° C, protected from light. Once opened, use the product within 7 days. Discard the unused product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Bottle containing 100 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No. 133

DIARREVET® ORAL POWDER

POLVO ORAL.

ANTIDIARREICO.

Technical Specification

SPECIES

Bovine, horses, pigs, ovine and goats.

Control diarrhea caused by germs sensitive to sulfonamides (*Salmonella spp, E.coli* and *Coccidia*).

DOSAGE FORM

Oral powder.

THERAPEUTIC ACTION

Anti-diarrheal.

COMPOSITION

Each 100 g contains:
Sulfaguanidine Monohydrate ..... 15 g
Phthalylsulfathiazole ..... 5 g
Citrus Pectin ..... 1 g
Bentonite ..... 2 g
Colloidal Kaolin ..... 77 g

INDICATIONS

Do not use in animals hypersensitive to sulfonamides.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route:

Dissolve the recommended dose in 100 to 200 mL of warm water.

Product dose: (1 tablespoon equal to 10 g of the product)

	Bovine and adult horses	Ovines, Goats, Calves and Fouls	Adult Pigs	Piglets
INITIAL DOSE	1 tablespoon (for every 500 Kg of weight: Sulfaguanidine 3g; Phthalysulfatiazole 1g; Pectin 0,2g; Bentonite 0,4g; Kaolin 15,4g)	1 tablespoon (for every 50 Kg of weight: Sulfaguanidine 1,5g; Phthalylsulfatiazole 0,5g; Pectin 0,1g; Bentonite 0,2g; Kaolin 7,7g)	1 ½ tablespoon (for every 100 Kg of weight: Sulfaguanidine 2,25 g; Phthalylsulfatiazole 0,75 g; Pectin 0,15 g; Bentonite 0,3 g; Kaolin 11,55 g)	½ tablespoon (for every 30 Kg of weight: Sulfaguanidine 0,75 g; Phthalylsulfatiazole 0,25 g; Pectin 0,05 g; Bentonite 0,1 g; Kaolin 3,85 g)
MAINTENANCE DOSE	1 tablespoon (for every 500 Kg of weight: Sulfaguanidine 1,5 g; Phthalylsulfatiazole 0,5 g; Pectin 0,1 g; Bentonite 0,2 g; Kaolin 7,7 g) every 8 hours for 4 days.	1 tablespoon)for every 50 Kg of weight: Sulfaguanidine 1,5 g; Phthalylsulfatiazole 0,5 g; Pectin 0,1 g; Bentonite 0,2 g; Kaolin 7,7 g) every 8 hours for 4 days.	1 tablespoon (for every 100 Kg of weight: Sulfaguanidine 1,5 g; Phthalylsulfatiazole 0,5 g; Pectin 0,1 g; Bentonite 0,2 g; Kaolin 7,7 g) every 8 hours for 4 days.	¼ tablespoon (for every 30 Kg of weight: Sulfaguanidine 0,375 g; Phthalylsulfatiazole 0,125 g; Pectin 0,025 g; Bentonite 0,05 g; Kaolin 1,925 g) every 8 hours for 4 days.

DRUG INTERACTIONS

Adsorbents such as Kaolin and Pectin can inhibit the intestinal absorption of other drugs that are administered together or during treatment with DIARREVET®.

CONTRAINDICATIONS

Do not use in animals hypersensitive to sulfonamides.

PRECAUTIONS

Use with caution in animals with kidney or liver failure.

SPECIAL PRECAUTIONS FOR USE

Use with caution in animals with kidney or liver failure.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Meat: 10 days (cattle, pigs, sheep and goats). Do not administer in horses whose meat is intended for human consumption.
Milk: 4 days (cattle). Do not administer in sheep or goats whose milk is intended for human consumption.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 ° and 30 ° C, protected from light. Use immediately once prepared and discard the excess product.

CONDITION OF SALE

Supply only on veterinary prescription.

PRESENTATION

Box with 25 sachets of 50 g each.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 942
- Dominican Republic: Reg. No. 5608

DIPRAMIDA® ORAL SOLUTION

SOLUCIÓN ORAL.

ANTIEMÉTICO - GASTROCINÉTICO.

Technical Specification

SPECIES

Dogs and cats.

Indicated in the management of vomiting of diverse etiology, in reflux esophagitis and in gastric motility disorders. It is also indicated in the preparation of radiological studies of the digestive tract.

DOSAGE FORM

Oral solution.

THERAPEUTIC ACTION

Antiemetic – Gastrokinetic.

COMPOSITION

Each 1 mL of oral solution contains:
Metoclopramide hydrochloride monohydrate .....5.27 mg
(Equivalent to 5 mg of Metoclopramide hydrochloride)
Excipients q.s.p .....1 mL

INDICATIONS

- Do not administer in cases of gastrointestinal obstruction or perforation.
- Do not administer in patients with hypersensitivity to Metoclopramide.
- Do not administer to pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration:

Active ingredient dose:
0.25 – 0.5 mg/Kg of weight, three times a day.

Product dose:
1 - 2 drops/Kg of weight, three times a day.

Administer 30-45 minutes before the meal and at the time of sleep.

DRUG INTERACTIONS

- Avoid concomitant administration with anticholinergics. Atropine may block the effect of Metoclopramide on gastrointestinal motility.
- Phenothiazine tranquilizers can potentiate the effects centrally.

CONTRAINDICATIONS

- Do not administer in cases of gastrointestinal obstruction or perforation.
- Do not administer in patients with hypersensitivity to Metoclopramide.
- Do not administer to pregnant or lactating females.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- In the case of contact with the skin or eyes, it is recommended to wash with plenty of water. If irritation develops and persists, see a doctor.
- In case of ingestion, do not induce vomiting. Get medical help.

WARNINGS

Advertencias y precauciones especiales de uso:

- Mantener fuera del alcance de los niños.
- Se recomienda administrar con precaución en pacientes epilépticos.

SIDE EFFECTS

ADVERSE EFFECTS

In dogs, the most common adverse reactions, although rare, include changes in mental status and behavior (restlessness, involuntary spasms, aggression, hyperactivity or drowsiness, and depression). Cats may show signs of excitement or disorientation. Both species can develop constipation in prolonged treatments.

OBSERVATIONS

Shake before use.

Special precautions for the disposal of unused product or waste material:
Empty containers can be discarded as household waste, without any special precautions. Do not dispose of containers with product residues on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 ° and 30 ° C, protected from light. Once the container is opened, use within 12 weeks. Discard the unused product after that period of time.

CONDITION OF SALE

Supply only on veterinary prescription.

PRESENTATION

Bottle containing 20 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG N°: 1818
Perú: Registro SENASA F.43.32.I.0020

COUNTRIES WHERE IT IS MARKETED

Imported and Distributed in Peru by:

Representaciones Durand SAC.
Av. Manuel Olguín N° 501 Oficina N° 604
Santiago de Surco Lima.

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

DOGUIVIT® ADULT - TABLETS

COMPRIMIDO ORAL.

MULTIVITAMÍNICO Y MINERALES PARA PERROS ADULTOS.

Technical Specification

SPECIES

Dogs.

It is indicated in animals with deficiencies of the vitamins and minerals formulated, in the growth, pregnancy, lactation, stress or convalescence.

DOSAGE FORM

Tablets.

THERAPEUTIC ACTION

Multivitamin and minerals for adult dogs.

COMPOSITION

Each tablet contains:

Iron sulfate anhydrous.....25.84 mg

(Equivalent to 5.2 mg of Iron (base))

Cyanocobalamin 1% FG.....0.80 mg

(Equivalent to 0.008 mg of Vitamin B12)

Folic Acid.....0.2 mg

Riboflavin 5-phosphate, sodium salt.....1.37 mg

(Equivalent to 1.0 mg of Riboflavin (base))

Nicotinamide.....10.0 mg

Retinyl Acetate (Vitamin A).....1250 IU

Cholecalciferol (Vitamin D3).....125 IU

dl-alpha-Tocopheryl Acetate (Vitamin E).....2 IU

Pyridoxine Hydrochloride (Vitamin B6).....1.0 mg

Thiamine Hydrochloride (Vitamin B1).....1.0 mg

Calcium Panthotenate.....0.5 mg

Calcium Hydrogen Phosphate.....258.0 mg

(Equivalent to 60 mg of Calcium and 46.5 mg of Phosphorus)

Magnesium Sulfate heptahydrate.....10.23 mg

(Equivalent to 1.0 mg of Magnesium)

Excipients q.s.p.....1 tablet

ROUTE OD ADMINISTRATION AND DOSAGE

Give directly or crushed together with food.

Suggested daily dose:

- Dogs of 5 kg or less: 1 tablet
- Dogs over 5 kg: 2 tablets

Administer for at least 1 month

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a fresh, dry place, out of direct light, at no more than 30°C.

CONDITION OF SALE

OTC product (non-prescription)

PRESENTATION

30 tablets.

PREPARED BY

Drag Pharma Laboratory.

DOGUIVIT® PUPPY - TABLETS

COMPRIMIDO ORAL.

MULTIVITAMÍNICO Y MINERALES PARA CACHORROS.

Technical Specification

SPECIES

Puppies.

Multivitamin for puppies. Chewable tablets with pleasant taste for dogs under 1 year of age.

DOSAGE FORM

Tablets.

THERAPEUTIC ACTION

Multivitamin and minerals for puppies.

COMPOSITION

Each tablet contains:

DL- Methionine.....10 mg

Lysine Hydrochloride.....2.5 mg

Calcium Pantothenate.....0.45 mg

Nicotinamide.....1.8 mg

Riboflavin-5-Phosphate.....0.45 mg

Vitamin B1.....0.15 mg

Vitamin B6.....0.15 mg

Vitamin B12.....0.1 µg

Vitamin A.....500 IU

Vitamin D.....50 IU

Vitamin E.....1 IU

Calcium.....138 mg

Phosphorus.....108 mg

Magnesium.....5 mg

Iodine.....0.025 mg

Zinc.....0.05 mg

Copper.....0.1 mg

Potassium.....25 mg

Sodium.....25 mg

Iron.....1 mg

Manganese.....0.5 mg

Excipients q.s.p.....1 tablet

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route; give directly or crushed together with food.

Suggested daily dose:

- Dogs of 5 kg or less: 2 tablets
- Dogs over 5 kg: 4 tablets
- For convalescent animals, increase the dose by 50% or as indicated by your veterinarian.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

IT IS NOT A COMPLETE FOOD.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

60 tablets

PREPARED BY

Drag Pharma Laboratory.

RECORDS

SAG Inscription #: RM03-010

DOGUIVIT® SENIOR - TABLETS

COMPRIMIDO ORAL.

MULTIVITAMÍNICO Y MINERALES PARA PERROS SENIOR.

Technical Specification

SPECIES

Dogs.

Multivitamin for aged dogs.

DOSAGE FORM

Tablets.

THERAPEUTIC ACTION

Multivitamin and minerals for aged dogs.

COMPOSITION

Each tablet contains:

Vitamin A.....1.250 IU

Vitamin D3.....125 IU

Vitamin E.....2 IU

Riboflavin 5 Phosphate, Sodium.....1.37 mg

Vitamin B1.....1.0 mg

Vitamin B6.....1.0 mg

Vitamin B12.....8 µg

Folic Acid.....0.2 mg

Dried extract of Schizandra.....30 mg

Chondroitin Sulfate.....100 mg

Calcium Panthotenate.....0.5 mg

Nicotinamide.....10 mg

Calcium.....60 mg

Phosphorus.....46.5 mg

Magnesium.....1 mg

Iron (II) sulfate.....25.84 mg

Excipients q.s.p.....1 tablet

ROUTE OD ADMINISTRATION AND DOSAGE

Oral. Give directly or crushed together with food.

Suggested daily dose:

- Dogs of 5 kg or less: 1 tablet
- Dogs over 5 kg: 2 tablets
- For convalescent animals, increase the dose by 50% or as indicated by your veterinarian.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

IT IS NOT A COMPLETE FOOD.

CONSERVATION

Store in a fresh, dry place, out of direct light, at no more than 30°C.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

30 tablets.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- SAG Inscription #: RM03-010
- Panama: Reg. No. RF-4243-09

DORACTINA® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANTIPARASITARIO DE EFECTO ENDECTOCIDA DE AMPLIO ESPECTRO Y ACCIÓN PROLONGADA.

Technical Specification

SPECIES

Bovine.

For the treatment of parasitic infections caused by gastrointestinal and pulmonary nematodes and external parasites such as sucking lice and scabies mites in cattle.

Doramectin is a broad spectrum anti-parasitic and long lasting effect because of its lipophilic properties. It is effective against adult and larvae parasites from the following species:

- **Nematodes:** *Ostertagia spp., Cooperia spp., Haemonchus spp., Trichostrongylus spp.,Oesophagostomum spp., Nematodirus spathiger, Bunostomum spp., Strongyloides spp., Trichuris spp., Dictyocaulus viviparus.*
- **Lice:** *Haematopinus eurysternus, Linognathus vituli, Solenoptes capillatus.*
- **Mites:** *Psoroptes bovis, Sarcoptes scabiei, var. bovis.*

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Endectocide antiparasitic of broad spectrum and long-acting effect.

COMPOSITION

Each 100 mL of product contains:
Doramectin.....1.0 g
Excipients q.s.p.....100 mL

PROPERTIES

Doramectin belongs to the family of Avermectins and presents a similar structure to Ivermectin. It has a broad spectrum of anti-parasitic activity, causing paralysis of the nematodes and arthropods. The main mechanism of action of the active component in Doractina (Doramectin) is to inhibit the electrical activity controlling nerve cells in nematodes and muscle cells in arthropods, causing paralysis and death of the parasite. This occurs because de Doramectin joins the chloride channels linked to glutamate in the nerves (nematodes) and muscle cells (arthropods). This reaction results in an increase of the cell membrane permeability to chloride ions, causing hyperpolarization of the affected cells and subsequently paralysis and death of the parasite. Doramectin can also join to the chloride ion channels linked to GABA (Gamma-monobutyric acid) causing the same effect. Doramectin is widely distributed in the body and as a lipophilic substance is concentrated in the adipose tissue. This leads to an extended resistance in plasma because of its slow release time. Because of its lipophilic characteristics, the maximum doramectine concentration in bovine plasma is reached at 3 hours after the subcutaneous administration.

INDICATIONS

- Do not use in unauthorized species.
- In dogs, especially the Collie breed and its crossbreeds, like other avermectins, it can cross the blood-brain barrier with serious consequences.

ROUTE OD ADMINISTRATION AND DOSAGE

Subcutaneous route.

Active ingredient dose:
200 µg/Kg of weight, single dose.

Product dose:

1 mL for every 50 Kg of weight, single dose.

For collective treatment, it is recommended to use an automatic dosing syringe.

CONTRAINDICATIONS

- Do not use in unauthorized species.
- In dogs, especially the Collie breed and its crossbreeds, like other avermectins, it can cross the blood-brain barrier with serious consequences.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Meat: 42 days.

Milk: Do not use in animals whose milk is intended for human consumption.

CONSERVATION

Store in a cool and dry place between 2° and 30°C and protected from sunlight.

CONDITION OF SALE

Supply only on veterinary prescription.

PRESENTATION

Ampule with 50 mL and 250 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG N° 2064-B
Bolivia: Reg. SENASAG PUV-F N° 007254/16
Perú: Registro SENASA F.54.01.I.0200

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:
AGROGUARANI SRL
TEL: + (591) 314-1401
Santa Cruz de la Sierra, Bolivia.

Imported and Distributed in Peru by:
Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

DORAZEL® PLUS - SHAMPOO

SHAMPOO.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Dogs.

Effective in eliminating fleas (*Ctenocephalides felis* and *Ctenocephalides canis*), ticks (*Rhipicephalus sanguineus*) and lice (*Trichodectes canis*) in dogs.

DOSAGE FORM

Shampoo.

THERAPEUTIC ACTION

External antiparasitic.

COMPOSITION

Each 100 mL of shampoo contains:
Permethrin.....1 g
Piperonyl Butoxide.....2.5 g
Excipients q.s.p.....100 mL

INDICATIONS

- Do not apply in cats.
- Do not administer in animals with hypersensitivity to any of the active principles.
- Do not administer to puppies under 12 weeks of age.
- Do not administer in pregnant or lactating females.

MODE OF APPLICATION

- Shake the container before using, then wet the animal with plenty of water and apply a sufficient amount of shampoo from the head to tail, directly on the skin and coat.
- Rub and let act for 10 to 15 minutes, then rinse with plenty of water.
- Repeat the treatment weekly, if necessary.

ROUTE OD ADMINISTRATION AND DOSAGE

For external use.

CONTRAINDICATIONS

- Do not apply in cats.
- Do not administer in animals with hypersensitivity to any of the active principles.
- Do not administer to puppies under 12 weeks of age.
- Do not administer in pregnant or lactating females.

PRECAUTIONS

- Avoid contact with your pet's eyes, mouth or other mucous membranes.
- In case of irritation, discontinue use of the product and wash with plenty of water.

SPECIAL PRECAUTIONS FOR USE

- Avoid contact with your pet's eyes, mouth or other mucous membranes.
- In case of irritation, discontinue use of the product and wash with plenty of water.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- The use of gloves is recommended at the time of application.
- Wash your hands with plenty of water after use.
- In case of contact with the eyes, mouth or skin, wash the affected area with plenty of water.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Do not throw the empty container or with product remains, in rivers, lakes or streams of natural water. Dispose of the waste of this product with care together with household waste.

CONSERVATION

Keep in a fresh, dry place, at room temperature, between 15 and 30 °C.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

300 mL bottle

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. S.A.G. No.: 0402

DORAZEL® SHAMPOO

SHAMPOO.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Dogs.

Antiparasitic shampoo, effective to eliminate fleas (*Ctenocephalides canis* and *Ctenocephalides felis*), ticks (*Rhipicephalus sanguineus*) and lice (*Trichodectes canis*).

DOSAGE FORM

Shampoo.

THERAPEUTIC ACTION

Antiparasitic shampoo.

COMPOSITION

Each 100 mL of shampoo contains:
Permethrin.....1 g
Excipients q.s.....100 mL

INDICATIONS

Do not use in puppies under 12 weeks of age, pregnant and lactating females or in animals with hypersensitivity to Permethrin.

MODE OF APPLICATION

1. Wet your pet using warm water, then apply a sufficient quantity of shampoo on the whole body surface. Rub for 2 to 3 minutes until obtaining abundant lather.
2. Let act for 10 to 15 minutes, and then rinse.
3. Repeat the procedure, if necessary.

ROUTE OD ADMINISTRATION AND DOSAGE

For external use.

CONTRAINDICATIONS

Do not use in puppies under 12 weeks of age, pregnant and lactating females or in animals with hypersensitivity to Permethrin.

PRECAUTIONS

- Avoid contact with the eyes, mouth or other mucous membranes.
- In case of irritation, discontinue use of the product and wash with plenty of water.
- Do not throw the container into water courses.
- Dispose of remains of the product together with household waste.

SPECIAL PRECAUTIONS FOR USE

- Avoid contact with the eyes, mouth or other mucous membranes.
- In case of irritation, discontinue use of the product and wash with plenty of water.
- Do not throw the container into water courses.
- Dispose of remains of the product together with household waste.

SPECIAL PRECAUTIONS FOR THE OPERATOR

The use of gloves is recommended at the time of application.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Storage in a fresh, dry place, at room temperature, between 15 and 30 °C.
Once the container is opened, use within 12 weeks. Discard the unused product after that period.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

100 mL bottle

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. S.A.G. No.: 0392

DOXIMICIN® ORAL SOLUTION

SOLUCIÓN ORAL.

ANTIMICROBIANO DE AMPLIO ESPECTRO.

Technical Specification

SPECIES

Dogs and cats.

Doximicin® is indicated for the treatment of upper and lower respiratory tract infections caused by sensitive etiological agents, such as primary infectious pictures or those associated with respiratory virosis, bacterial skin infections, eye and peri-eye infections caused by Chlamydiae or other strains Sensitive, Canine Ehrlichiosis and Canine and Feline Mycoplasmosis (Hemobartonellosis).

Presentation in oral solution, in addition to facilitating administration, avoids the risk of causing esophageal narrowing in cats.

DOSAGE FORM

Oral solution.

THERAPEUTIC ACTION

Broad spectrum antimicrobial.

COMPOSITION

Each mL of the product contains:
Doxycycline Hyclate.....11.08 mg
(Equivalent to 10 mg of Doxycycline basic)
Excipients q.s.p.....1 mL

INDICATIONS

Do not use **Doximicin®** in animals hypersensitive to Doxycycline.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route.

- **Dogs and cats:** 10 mg/Kg, once a day for 14 to 21 days.
- **Product dose:** 1 mL per Kg of weight, once a day for 14 to 21 days. In respiratory tract infections, administer it for a maximum of 10 days.

CONTRAINDICATIONS

Do not use **Doximicin®** in animals hypersensitive to Doxycycline.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a dry place, at room temperature between 15° and 30°C and protected from sunlight.

CONDITION OF SALE

Supply only on veterinary prescription.

PRESENTATION

In a bottle with 60 ml. Includes dosing syringe and dropper.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No 2029-B

DOXIMICIN® ORAL TABLETS

COMPRIMIDO ORAL.

ANTIMICROBIANO DE AMPLIO ESPECTRO.

Technical Specification

SPECIES

Dogs and cats.

Broad-spectrum antimicrobial indicated for the treatment of diseases caused by agents susceptible to Doxycycline such as: respiratory tract infections (upper and lower) caused by *Mycoplasma spp.*, *Chlamydia spp.*, *Bordetella spp.*, *Haemophilus spp.*, *Actinomyces spp.* and *Pasteurella multocida*; skin infections caused by *Streptococcus spp.* and *Staphylococcus spp.*, eye and peri-ocular infections caused by *Chlamydia spp.*; Canine *ehrlichiosis* caused by *Ehrlichia canis* and feline and canine infectious anemia caused by *Haemobartonella felis* and *Haemobartonella canis*, respectively.

DOSAGE FORM

Oral tablet.

THERAPEUTIC ACTION

Broad spectrum antimicrobial.

COMPOSITION

Each oral tablet contains:
Doxycycline Hyclate.....100 mg
(Equivalent to 86.6 mg de Doxycycline basic anhydrous)
Excipients q.s.p.....1 Tablet

INDICATIONS

- Do not administer to animals hypersensitive to tetracyclines such as Doxycycline.
- Do not administer to animals with dysphagia problems or vomiting.
- Do not administer during pregnancy, lactation, or in breeding animals.
- Do not administer to puppies under 7 - 8 weeks of age.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route.

- **Dose of the active principle:** 10 mg of Doxycycline/Kg of weight, every 24 hours for 14 to 21 days. In respiratory tract infections, administer for a maximum of 10 days.
- **Product dose:** Administer 1 tablet for every 10 Kg of weight, every 24 hours for 14 to 21 days. In respiratory tract infections, administer for a maximum of 10 days. It can be administered together with food.

DRUG INTERACTIONS

- Simultaneous administration of oral absorbents, iron preparations, antacids and, in general, gastrointestinal products containing aluminum, calcium, magnesium, zinc or bismuth cations reduces the bioavailability of doxycycline.
- The half-life of doxycycline is reduced when co-administered with barbiturates or phenytoin.
- There may be cross-resistance with other tetracyclines.
- Doxycycline should not be used in conjunction with other antibiotics such as β-lactams.

CONTRAINDICATIONS

- Do not administer to animals hypersensitive to tetracyclines such as Doxycycline.
- Do not administer to animals with dysphagia problems or vomiting.
- Do not administer during pregnancy, lactation, or in breeding animals.
- Do not administer to puppies under 7 - 8 weeks of age.

PRECAUTIONS

-Keep out of the reach of children and pets.
-Do not exceed the recommended doses.

SPECIAL PRECAUTIONS FOR USE

-Keep out of the reach of children and pets.
-Do not exceed the recommended doses.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not handle by people hypersensitive to doxycycline.
- Wash your hands with plenty of water immediately after handling the product.
- In case of accidental ingestion, go immediately to a medical center and show the product insert.

WARNINGS

- Mantener fuera del alcance de los niños.
- No exceder las dosis recomendadas.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be discarded as domestic waste, without any special precautions. Do not dispose of containers with product residues on the ground or in watercourses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 °C, protected from light.

CONDITION OF SALE

Supply only on veterinary prescription.

PRESENTATION

Box containing 10 tablets.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG N° 1320-B
Costa Rica: Reg. N° MV-5869
Bolivia: Reg. SENASAG PUV-F N° 007261/16

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

DRAGBUTEROL® ORAL GEL

GEL ORAL.

BRONCODILATADOR.

Technical Specification

SPECIES

Horses.

Dragbuterol® is mainly recommended for the treatment of obstructive diseases of the respiratory tract in horses, such as chronic obstructive pulmonary disease and allergic respiratory diseases. Because of its mucolytic action, it is also recommended as adjuvant in the treatment of infectious diseases such as pneumonia and bronchitis.

DOSAGE FORM

Oral gel.

THERAPEUTIC ACTION

Bronchodilator.

COMPOSITION

Each 100 mL of gel contains:
Clenbuterol Hydrochloride..... 2.0 mg
Excipients q.s.p.....100 mL

INDICATIONS

- Do not administer in animals hypersensitive to Clenbuterol Hydrochloride.
- Do not administer in horses with cardiovascular disorders.
- Do not administer in mares in advanced gestation or in lactation period.
- Do not administer to breeding males.
- Do not administer to offspring.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route.

Dose of the active principle:
- Initial dose: 0.8 µg of Clenbuterol Hydrochloride / Kg of b.w. every 12 hours.

If after 3 days of treatment (6 doses) no improvement is observed, increase the dose according to the following scheme:
- 1.6 µg of Clenbuterol Hydrochloride / Kg of b.w. every 12 hours for 3 days, if there is no improvement,
- 2.4 µg of Clenbuterol Hydrochloride / Kg of b.w. every 12 hours for 3 days, if there is no improvement,
- 3.2 µg of Clenbuterol Hydrochloride / Kg of b.w. every 12 hours for 3 days.

Product dosage:
- Initial dose and mild obstructive symptoms: 4 mL / 100 Kg of b.w., every 12 hours for 30 days.
- Severe broncho-obstructive symptoms: maximum dose of 12-16 mL / 100 Kg of b.w. every 12 hours for 30 days.

If after 3 days of treatment (6 doses) with the initial dose no improvement is observed, increase the dose according to the following scheme:
- 8 mL / 100 Kg of b.w., every 12 hours for 3 days, if there is no improvement,
- 12 mL / 100 Kg b.w., every 12 hours for 3 days, if there is no improvement,
- 16 mL / 100 Kg of b.w., every 12 hours for 3 days.

Once the therapeutic dose is established, continue the treatment for 30 days. If the signs reappear, the treatment can be repeated for a further 30 days, but the determination of the effective dose must be done again in stages.
If after administering the higher dose no effect is observed, it means that the animal is non-responsive to Clenbuterol or that the obstructive condition is not of bronchial origin, then the treatment should be suspended.

DRUG INTERACTIONS

- Concomitant administration with other sympathomimetic amines (for example Terbutaline) can increase the adverse effects of Clenbuterol.
- The use with inhalation anesthetics (eg Halothane, Isofluorane) can predispose to ventricular arrhythmias, particularly in patients with pre-existing heart disease.
- The use with digitalis glycosides may increase the risk of cardiac arrhythmias.
- β-blockers (for example Propanolol) can antagonize the effects of Clenbuterol.
- Clenbuterol can antagonize the effects of Prostaglandin F2 and Oxytocin.

CONTRAINDICATIONS

- Do not administer in animals hypersensitive to Clenbuterol Hydrochloride.
- Do not administer in horses with cardiovascular disorders.
- Do not administer in mares in advanced gestation or in lactation period.
- Do not administer to breeding males.
- Do not administer to offspring.

PRECAUTIONS

- It is not recommended to start treatment with a dose higher than the established starting dose.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR USE

- It is not recommended to start treatment with a dose higher than the established starting dose.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not handle by people who are hypersensitive to Clenbuterol Hydrochloride.
- Do not handle by pregnant women.

SIDE EFFECTS

ADVERSE EFFECTS

Adverse effects and adverse reactions:
Moderate tachycardia, sweating, muscle tremors, hives, and restlessness may occur during the first days of treatment. The presence of ataxia is also described, although in low frequency.

GUARD PERIOD

Do not administer in animals whose meat is intended for human consumption.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 °C, protected from light. Once the container is opened, use within 3 days. Discard unused product after that time period.

CONDITION OF SALE

Sale with retained Veterinary Medical prescription only.

PRESENTATION

50 mL bottle

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No 1866

ECO-CAT® COLLAR

COLLAR.

COLLAR REPELENTE DE PULGAS PARA GATOS.

Technical Specification

SPECIES

Cats.

This necklace contains natural flea repellent essences. These oils have been selected for their effectiveness, their good tolerance on the skin and their pleasant lemon scent. Permanent use of the collar helps prevent flea infestation. This product is impervious to water and does not disturb your cat's sense of smell. An anti-strangulation safety device breaks the collar in the event of violent jerks.
The use of this collar associated with proper flea control in your pet's environment is recommended.

DOSAGE FORM

Collar.

THERAPEUTIC ACTION

Flea repellent collar for cats.

COMPOSITION

Each collar contains:
Lemon essential oil 7.4% (limonene, citrones, terpenes)
Inert substances q.s.p.....1 collar

PROPERTIES

- With natural oils
- Does not contain insecticidal substances
- Long acting
- With anti-choke device

USE INSTRUCTIONS

- Pull out and unroll the collar and then insert the end into the buckle.
- Fit the collar loosely around the dog's neck, and fix the free part of the collar to the clip.
- It is recommended to leave a space of two fingers between the collar and the neck.

WARNINGS

- Mantener fuera del alcance de los niños.
- Consulte a su Médico veterinario antes de usar este producto en animales debilitados, envejecidos, hembras preñadas, en amamantamiento, o animales bajo medicación.
- Suspender el uso si se desarrolla cualquier efecto no deseado.

OBSERVATIONS

For external use.

CONSERVATION

Store in a fresh, dry place, out of direct light, at no more than 30°C

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

1 collar.

PREPARED BY

Societé AB7 Industries S.A. Chemin des Monges Deyme, France.
Imported and distributed in Chile by Drag Pharma Laboratory.

ECO-DOG® COLLAR

COLLAR.

COLLAR REPELENTE DE PULGAS PARA PERROS.

Technical Specification

SPECIES

Dogs

This necklace contains natural flea repellent essences. These oils have been selected by its effectiveness, its good tolerance on the skin and its pleasant lemon scent. Permanent use of the collar Helps prevent flea infestation. This product is waterproof and does not disturb the sense of your dog's smell. An anti-strangulation safety device breaks the collar in case of violent jerks.

The use of this collar is recommended associated with proper management in the control of fleas in the environment of your pet.

DOSAGE FORM

Collar.

THERAPEUTIC ACTION

Flea repellent collar for dogs.

COMPOSITION

Each collar contains:
Lemon essential oil 7.4% (limonene, citrones, terpenes)
Inert substances q.s.p.....1 collar

PROPERTIES

- With natural oils
- Does not contain insecticidal substances
- Long acting
- With anti-choke device

USE INSTRUCTIONS

- Pull out and unroll the collar and then insert the end into the buckle.
- Fit the collar loosely around the dog's neck, and fix the free part of the collar to the clip.
- It is recommended to leave a space of two fingers between the collar and the neck.

WARNINGS

- Mantener fuera del alcance de los niños.
- Consulte a su Médico veterinario antes de usar este producto en animales debilitados, envejecidos, hembras preñadas, en amamantamiento, o animales bajo medicación.
- Suspender el uso si se desarrolla cualquier efecto no deseado.

OBSERVATIONS

For external use.

CONSERVATION

Store in a fresh, dry place, out of direct light, at no more than 30°C

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

1 collar.

PREPARED BY

Societé AB7 Industries S.A. Chemin des Monges Deyme, France.
Imported and distributed in Chile by Drag Pharma Laboratory.

ELECTROVET® ORAL SOLUTION

SOLUCIÓN ORAL.

SOLUCIÓN ORAL ISOTÓNICA REHIDRATANTE PARA PERROS Y GATOS, LISTA PARA BEBER.

Technical Specification

SPECIES

Dogs and cats.

Electrovet® is a pleasant tasting isotonic oral solution for dogs and cats based on glucose, glycine and electrolytes that helps in pet rehydration therapy. The presence of the amino acid glycine in the formulation, together with glucose, increases the intestinal absorption of sodium and water, helping to restore the deficit of water and electrolytes, which favors the rehydration of the pet after intense exercise, prolonged periods without ingestion of water, periods of convalescence, trips or fluid losses.

DOSAGE FORM

Oral solution

THERAPEUTIC ACTION

Isotonic Oral Rehydrating Solution for Dogs and Cats, ready to drink.

COMPOSITION

Each 100 mL of product contains:
Glucose ..... 2.23 g
Glycine ..... 0.31 g
Sodium Chloride ..... 0.43 g
Monobasic Potassium Phosphate ..... 0.20 g
Citric Acid Monohydrate ..... 0.03 g
Potassium Citrate ..... 0.01 g
Excipients q.s.p ..... 100 mL

INGREDIENTS

Purified Water, Glucose, Glycine, Sodium Chloride, Monobasic Potassium Phosphate, Citric Acid Monohydrate, Potassium Citrate, Sodium Benzoate, Dyes and Flavors allowed.

ROUTE OD ADMINISTRATION AND DOSAGE

Dosage and administration:
Administer orally through the dosing syringe or directly on the drinking plate for voluntary consumption by the pet.

- **Puppies and kittens:** 40-50 mL / day.
- **Cats:** 125-250 mL / day.
- **Dogs up to 5 Kg:** 125-250 mL / day.
- **Dogs up to 10 Kg:** 500 mL / day.
- **Dogs over 10 Kg:** up to 750 mL / day.

The dose should be divided into two to three administrations daily. Administer as needed, for a period of two days.

WARNINGS

- Mantener fuera del alcance de los niños
- En caso que su mascota se encuentre deshidratada debe ser evaluada por un Médico Veterinario.

OBSERVATIONS

- Does not constitute a complete food
- Exclusive use in animal feed

CONSERVATION

- Store protected from light below 30ºC.
- Once opened, consume within a week, keep refrigerated.

CONDITION OF SALE

OTC con prescription

PRESENTATION

bottle with 500 mL

PREPARED BY

Laboratorio Drag Pharma.

RECORDS

Chile: Reg. LENAA N°: RM 03-008N
Bolivia: Reg. SENASAG PUV-A nº 008926/19
Costa Rica: Lic. DAA-MAG 579-016

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

Imported and distributed in Costa Rica by:
Proventas de Cartago S.R.L.
100 meters east Hogares Crea, San Blas. Carthage.
Tel: 2591 4624 Fax: 2591 5339

LEVEL

1

ENDOFACIOL® BOVINE - ORAL SUSPENSION

SUSPENSIÓN ORAL.

ANTIPARASITARIO INTERNO DE AMPLIO ESPECTRO EN DOSIS ÚNICA
FASCIOLICIDA ORAL

Technical Specification

SPECIES

Bovine.

Indicated in the treatment and control of *Fasciola hepatic*a in its immature, juvenile and adult state; gastrointestinal and lung parasites, in their adult and larval stages in bovines. In ruminants it is active against inhibited larvae of *Ostertagia spp*.

DOSAGE FORM

Oral suspension.

THERAPEUTIC ACTION

Broad spectrum internal antiparasitic
Oral fasciolicide

COMPOSITION

Each 100 mL of suspension contains:
Ivermectin.....0.2 g
Triclabendazole.....12 g
Excipients q.s.p.....100 mL

INDICATIONS

Do not administer to animals with known hypersensitivity to ivermectins.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer orally, using a syringe or dosing gun.

Active ingredients dose:

- Ivermectin: 0.2 mg/Kg of weight.
- Triclabendazole: 12 mg/Kg of weight.

Product dose:
10 mL for each 100 kg of weight.

DRUG INTERACTIONS

Do not use concomitantly with other pharmaceuticals.

CONTRAINDICATIONS

Do not administer to animals with known hypersensitivity to ivermectins.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Aviod contact of the product with the skin and eyes. In case of contact, wash inmediately with plenty of water. If irritation exists and persists, see a doctor. Wash your hands after handling the product.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Meat: 49 days.

Do not administer to cows which milk is intended for human consumption.

OBSERVATIONS

Shake before use.

Special precautions for the disposal of unused product or waste material:
Dispose of this product with caution with household waste. Do not dispose of containers with product residues on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 ° and 30 ° C, protected from light. Once the container is opened, use within 12 weeks. Discard the unused product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

100 mL, 250 mL, 1 L and 2 L bottles.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG No. 1712-B
Rep. Dominicana: Reg. Nº 7724

ENDOFACIOL® EQUINE - ORAL SUSPENSION

SUSPENSIÓN ORAL.

ANTIPARASITARIO INTERNO DE AMPLIO ESPECTRO EN DOSIS ÚNICA Y FASCIOLICIDA.

Technical Specification

SPECIES

Horses.

Indicated in the treatment and control of *Fasciola hepatica* in its immature, juvenile and adult state; gastrointestinal and lung parasites, including migratory larvae; and larval stages of gastrophils in horses.

DOSAGE FORM

Oral suspension

THERAPEUTIC ACTION

Single dose broad spectrum internal antiparasitic and fasciolicide.

COMPOSITION

Each 100 mL of suspension contains:
Ivermectin.....0.2 g
Triclabendazole.....12 g
Excipients q.s.p.....100 mL

INDICATIONS

Do not administer to animals with known hypersensitivity to ivermectins.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer orally, using a syringe or a dosing gun.

Active ingredients dose:

- Ivermectin: 0.2 mg/Kg of weight.
- Triclabendazole: 12.0 mg/Kg of weight.

Product dose:
10 mL for each 100 kg of weight, single dose.

CONTRAINDICATIONS

Do not administer to animals with known hypersensitivity to ivermectins.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Aviod contact of the product with the skin and eyes. In case of contact, wash inmediately with plenty of water. If irritation exists and persists, see a doctor. Wash your hands after handling the product.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Do not administer to animals which meat is intended for human consumption.

OBSERVATIONS

Shake before use.

Special precautions for the disposal of unused product or waste material:
Dispose of this product with caution with household waste. Do not dispose of containers with product residues on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 ° and 30 ° C, protected from light. Once the container is opened, use within 12 weeks. Discard the unused product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

100 mL and 1 L bottles.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No. 1712-B

EQ-CASCO® ORAL POWDER

POLVO ORAL.

SUMPLEMENTO NUTRICIONAL QUE POTENCIA LA ESTRUCTURA DE CASCOS EN CABALLOS

Technical Specification

SPECIES

Horses.

EQ–Casco® regenerates through diet, frayed or cracked hooves. EQ-Casco® components will be improving the hoof and skin quality of your horse.

DESCRIPTION

Nutritional supplement that enhances the structure of hooves in horses

DOSAGE FORM

Oral powder.

THERAPEUTIC ACTION

Nutritional additive that strengthens the hoof structure.

COMPOSITION

Each 100 g contains:
Methionine.....10 g
Lysine HCl.....12 g
Zinc.....700 mg
Magnesium.....200 mg
Biotin 100%.....25 mg
Vitamin B6.....40 mg
Calcium Pantothenate.....60 mg
Oligosaccharides q.s.p.....100 g

OLIGOSACCHÁRIDES from *Saccharomyces cerevisiae*, which is rich in amino acids such: Cysteine, Tryptophan, Threonine, Isoleucine, Histidine, Valine, Leucine, Arginine, Phenylalanine; and vitamins such: Thiamine, Riboflavin, Folic acid.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration.

30 grams a day (one measure).

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

EXCLUSIVE USE IN ANIMAL FEED
DOES NOT CORRESPOND TO A COMPLETE FOOD

CONSERVATION

Store in a cool, dry place and protected from light, at no more than 30ºC

CONDITION OF SALE

Over the counter.

PRESENTATION

1 Kg container.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. LENAA N°: RM 03-008N

EQUIFLEX® NEXT LEVEL - SYRUP

JARABE.

SUPLEMENTO NUTRICIONAL CONDROPROTECTOR.

Technical Specification

SPECIES

Horses.

Equiflex Next Level is a nutritional supplement for horses. By its components, helps the maintenance of a healthy joint function and promotes the recovery of the cartilaginous matrix in injured joints.

DOSAGE FORM

Syrup.

THERAPEUTIC ACTION

Chondroprotector nutritional supplement.

COMPOSITION

Qualitative analysis:

Chondroprotectors:

Methylsulfonylmethane (MSM).....10%

Glucosamine Sulfate.....10%

Chondroitin Sulfate.....7%

Vitamin:

Vitamin C.....3.33%

Amino acid:

DL-Methionine.....1%

ROUTE OD ADMINISTRATION AND DOSAGE

500 Kg of weight, it recommends administering orally, 60ml for 5 days, then continue with 30ml daily.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

This is not a complete food.

CONSERVATION

Store in a cool and dry place between 15° and 30°C.

CONDITION OF SALE

Over the counter.

PRESENTATION

1 L bottle.

PREPARED BY

Drag Pharma Laboratory.

EQUIFLEX® SYRUP

JARABE.

SUPLEMENTO NUTRICIONAL.

Technical Specification

SPECIES

Horses.

This is an enriched nutritional supplement for horses. Intended to be used especially in animals subjected to continuous exercise.

DOSAGE FORM

Syrup.

THERAPEUTIC ACTION

Supplement Facts.

COMPOSITION

Equine supplement composition:

- Chondroprotectors: Chondroitin Sulfate and Glucosamine Sulfate.
- Amino-acids: Methionine.
- Vitamins: Thiamine, Pyridoxine and vitamin C.
- Minerals: Copper, Zinc, Manganese.

Qualitative analysis:
Glucosamine Sulfate min.....10%
Chondroitin Sulfate min..... 7%
Vitamin C min.....3.5%
Vitamin B1 min.....0.1%
Vitamin B6.....0.033%
Methionine min.....1.0%
Copper min.....0.085
Manganese min.....0.054%
Zinc min.....0.087%
Yucca extract 50%.....0.333%

ROUTE OD ADMINISTRATION AND DOSAGE

500 Kg horses, it recommends administering orally, 60ml for 5 days, then continue with 30 ml daily.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

This is not a complete food.

CONDITION OF SALE

Over the counter.

PRESENTATION

1 L bottle.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

EQUIFORT® ORAL POWDER.

POLVO ORAL.

SUPLEMENTO NUTRICIONAL CON VITAMINAS, MINERALES, AMINOÁCIDOS Y ENERGIZANTES.

Technical Specification

SPECIES

Horses.

Nutritional supplement with vitamins, minerals, amino acids and invigorating, especially formulated to support dietary requirement of pure blood running horses or horses in general.

DOSAGE FORM

Oral powder.

THERAPEUTIC ACTION

Nutritional supplement with vitamins, minerals, amino acids and invigorating, especially formulated to support dietary requirements of pure blood running horses and horses in general.

COMPOSITION

Each 100 g contains:

Sodium chloride ..... 50 g

Vitamin A.....200,000 U.I.

Vitamin D.....40,000 U.I.

Vitamin E.....100 U.I.

Vitamin K.....0.04 g

Vitamin B2.....0.2 g

Calcium Pantothenate .....0.15 g

Niacin F.G.....0.37 g

Vitamin B12.....0.0003 g

Vitamin B1.....0.15 g

VitaminaB6.....0.02 g

Folic acid .....0.015 g

Choline Hydrochloride .....0.45 g

Di-calcium Phosphate.....20 g

Calcium Gluconate.....2 g

Glucose.....2 g

Iodine.....0.0045%

Magnesium.....0.04%

Iron….....0.54%

Copper.....0.04%

Cobalt.....0.0045%

Zinc.....0.141%

Potassium.....0.0015%

Methionine….....0.1 g

Lysine.....0.2 g

Excipients q.s.p.....100 g

ROUTE OD ADMINISTRATION AND DOSAGE

- Adult horses: 25-30 g a day
- Training horses: 30-40 g a day.
- Pregnant mares: 30-50 g a day
- Foals: 20-30 g a day.
- Cattle: 25-30 g a day.

The dose is given mixed with the daily ration of food.

A dispenser is attached to manage the amount indicated.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

- Nice apple flavor.
- This is not a complete food.

CONDITION OF SALE

Over the counter.

PRESENTATION

1 Kg container.

PREPARED BY

Drag Pharma Laboratory.

EQUIFORT® SYRUP

JARABE.

SUPLEMENTO NUTRICIONAL MULTIVITAMÍNICO Y MINERAL.

Technical Specification

SPECIES

Horses.

Multi vitamin/mineral nutritional supplement for horses of any age, competition animals and horses with vitamin/mineral deficiency states.

DOSAGE FORM

Syrup.

THERAPEUTIC ACTION

Multi Vitamin/Mineral Nutritional Supplement.

COMPOSITION

Each 100 mL contains:

Copper.....150 mg

Cobalt .....6 mg

Potassium .....330 mg

Magnesium .....70 mg

Manganese .....130 mg

Zinc .....370 mg

Iron .....1.000 mg

Calcium Pantothenate .....170 mg

Folic acid .....35 mg

Biotin .....0.08 mg

Vitamin A .....80,000 U.I.

Vitamin D2 .....12,000 U.I.

Vitamin E .....150 U.I.

Vitamin B1 .....200 mg

Vitamin B2 .....100 mg

Vitamin B6 .....35 mg

Vitamin B12 .....400 µg

Sucrose.....50 g

Choline Chloride .....675 mg

Selenium .....3.1 mg

Excipients q.s.p. .....100 mL

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route or mixed with food.

- Foals: 30 mL/day for 3 to 5 days.
- Weak horses: 50 mL/day for 3 to 5 days.
- High demand horses: 100 mL/day for 3 to 5 days.
- Maintenance: 10 to 30 mL a day for 30 days.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

- With Apple flavor.
- Shake before use.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° and 30°C.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

500 mL, 1 and 3.8 L bottles.

PREPARED BY

Drag Pharma Laboratory.

EQUIGASTRIN® ORAL PASTE

PASTA ORAL.

INHIBIDOR DE LA SECRECIÓN GÁSTRICA.

Technical Specification

SPECIES

Horses.

Indicated for the treatment and prevention of gastric ulcers in adult horses and foals.

DESCRIPTION

Indicated for the treatment and prevention of gastric ulcers in adult horses and foals.

DOSAGE FORM

Oral paste.

THERAPEUTIC ACTION

Gastric secretion inhibitor.

COMPOSITION

Each 100 g of oral paste contains:
Omeprazole .....37.0 g
Excipients q.s.p. .....100 g

INDICATIONS

Contraindications, Warnings and Special Precautions for use:

- Keep out of the reach of children
- Do not administer to horses with known hypersensitivity to Omeprazole.
- Do not administer to pregnant or lactating females.
- Not recommended for horses less than 4 weeks old or with a body weight less than 70 kg.

MODE OF APPLICATION

- To administer the treatment dose, place the notch of the dispenser in the division corresponding to the body weight of the animal. The total content of the syringe will treat 1,000 Kg of weight at the rate of 4 mg of omeprazole /Kg of weight.

- To administer the maintenance or preventive dose, place the notch of the dispenser in the division corresponding to half the body weight of the animal. In this case the total content of the syringe will allow treatment of four 500kg animals at a rate of 2 mg of omeprazole /Kg de peso, or dose a 500 kg animal for four days.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration way: Oral.

For the treatment of gastric ulcers:
Administer a dose of 4 mg / Kg of body weight, once a day, for 28 consecutive days, followed by an administration of 2 mg / Kg of weight, once a day for 28 days, to reduce the recurrence of gastric ulcers after the treatment. If there is a recurrence, it is recommended to treat again with 4 mg / Kg of weight for 28 days.

For the prevention of gastric ulcers:
Administer a dose of 2 mg / Kg of body weight, once a day, for 28 consecutive days.

Practical product dose for a 500 kg horse:

Treatment of gastric ulcers:

- **From day 1 to 28**, administer ½ syringe, 1 time a day.
- **From the 29th to the 56th**, administer ¼ of a syringe, 1 time a day.

Recurrence (for 28 days): administer ½ syringe 1 time a day.

Prevention of gastric ulcers (for 28 days): administer ¼ syringe 1 time a day.

DRUG INTERACTIONS

- Omeprazole could decrease the hepatic clearance of Diazepam, Phenytoin, or Warfarin, so it is recommended not to apply together.
- Omeprazole can reduce the absorption of Ketoconazole, Ampicillin esters and Iron salts.

CONTRAINDICATIONS

Contraindications, Warnings and Special Precautions for use:

- Keep out of the reach of children
- Do not administer to horses with known hypersensitivity to Omeprazole.
- Do not administer to pregnant or lactating females.
- Not recommended for horses less than 4 weeks old or with a body weight less than 70 kg.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not eat, drink or smoke while handling the product
- Wash your hands after handling the product.
- Avoid contact of the product with the skin and eyes.

WARNINGS

ADVERSE EFFECTS

Unwanted effects and adverse reactions:
Omeprazole is generally well tolerated. However, gastrointestinal distress may occur (anorexia, colic, nausea, vomiting, flatulence, diarrhea).

GUARD PERIOD

Do not administer to horses which meat is intended for human consumption.

OBSERVATIONS

Precautions for Disposal of Unused Product or Waste Material:
Empty containers can be discarded as household waste, in their original container and tightly closed. Do not dispose of containers with product remains on the ground or water courses.
Contact the manufacturing company or companies specialized in waste disposal, to receive recommendations on the disposal of expired or unused products.

CONSERVATION

Store at a temperature between 2 and 30 ° C, protected from light. Once the container is opened, use within 8 days and keep between 15 and 30ºC, protected from light. Discard the unused product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

A box with 4 graduated syringes, with 10.82g each.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG No. 1954
Uruguay: Reg. MGAP N° 2017A00422

COUNTRIES WHERE IT IS MARKETED

Imported in Uruguay by:
VIVAFIL S.A. RIO NEGRO 1107 Montevideo - Uruguay, TEL 29001112
grupotecnovet@gmail.com
Technical Director: DMTV Diego Cuadrado.

EQUIMIC® ORAL GEL

GEL ORAL.

ANTIPARASITARIO INTERNO.

Technical Specification

SPECIES

Horses.

Single dose broad spectrum internal antiparasitic; effective against adult, migratory and / or larval stages of gastrointestinal parasites such as: *Strongylus vulgaris, Strongylus equinus, Strongylus edentatus, Oxyuris equi, Parascaris equorum*, small strongyles, *Strongyloides westeri, Trichostrongylus axei, Habronema spp. and Gasterophilus spp.*

DESCRIPTION

Broad spectrum internal antiparasitic for horses

DOSAGE FORM

Oral gel.

THERAPEUTIC ACTION

Internal Antiparasitic.

COMPOSITION

Each 1 mL of gel contains:
Ivermectin.....14.0 mg
Excipients q.s.p.....1 mL

INDICATIONS

- Do not use in foals under 4 weeks of age.
- Do not administer to horses with known hypersensitivity to Ivermectin.

MODE OF APPLICATION

- **Foals:** Start the treatment between 6-8 weeks of age, repeating the treatment at intervals of 6 to 8 weeks.
- **Mares:** It is recommended to treat 1 month before the due date and repeat 60 days after delivery.
- **Adult horses:** Treatment every 90 days. The repetition of the treatment depends on the parasite load and the season of the year.

ROUTE OD ADMINISTRATION AND DOSAGE

Route of administration: Oral.

Active ingredient dose:
0.2 mg/Kg of weight, single dose.

Product dose:
1 ml/70 kg of weight (0.2 mg/Kg), single dose.

Each syringe reaches 700 kg of weight.

CONTRAINDICATIONS

- Do not use in foals under 4 weeks of age.
- Do not administer to horses with known hypersensitivity to Ivermectin.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- In case of accidental ingestion, do not induce vomiting. Consult a doctor as soon as possible.
- In case of contact with skin or eyes, wash with plenty of water. If irritation occurs, consult a doctor.
- Wash hands after administration of the product.

WARNINGS

- Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

Adverse effects are observed in a low percentage of animals to which oral and injectable formulations containing Ivermectin are administered simultaneously.

In cases of severe infestations, signs of transient colic may be observed.

GUARD PERIOD

Do not administer to horses which meat is intended for human consumption.

OBSERVATIONS

Pleasant apple flavor.

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of water. Dispose of the waste of this product with care together with household waste.

CONSERVATION

Store in a cool, dry place, between 15 and 30 ° C.
Use immediately once opened and discard the excess product.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

10 mL graduated syringe.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG No. 1655-B

EQUIMIC® PLUS - ORAL PASTE

PASTA ORAL.

ANTIPARASITARIO DE AMPLIO ESPECTRO.

Technical Specification

SPECIES

Horses.

Equimic® Plus is a broad spectrum antiparasitic, because of the combination of Ivermectin and Praziquantel is effective against large strongyles (*Strongylus spp, Triodontophorus spp.*), small strongyles (*Cylicocyclus spp, Cylicostephanus spp, Cylicodontophorus spp, Gyalocephalus spp.*), pinworms (*Oxyuris equi*), ascarids (*Parascaris equorum*), trichostrongyles (*Trichostrongylus axei*), intestinal nematodes (*Strongyloides westeri*), stomach nematodes (*Habronema spp., Draschia megastoma*), horse bot fly (*Gasterophilus spp.*) and tapeworms (*Anoplocephala perfoliata*).

DOSAGE FORM

Oral paste.

THERAPEUTIC ACTION

Broad spectrum anti-parasite.

COMPOSITION

Each 20 g of oral paste contains:
Ivermectin.....0.18
Praziquantel.....1.35 g
Excipients q.s.p.....20 g

INDICATIONS

- Do not use in case of hypersensitivity to any of the active ingredients.
- Do not use in foals under 2 month old and stallions.

ROUTE OD ADMINISTRATION AND DOSAGE

Active ingredient dose:

- Ivermectin: 0.2 mg/Kg, single dose.
- Praziquantel: 1.5 mg/Kg, single dose.

Product dose:

3.3 g of Equimic® Plus for each 150 Kg of body weight, single dose. Each syringe is enough for up to 900 kg of weight.

Place the dosing Equimic® Plus syringe directly into the horse's mouth and download the volume equivalent to the recommended dosage according to the weight of the animal.

CONTRAINDICATIONS

- Do not use in case of hypersensitivity to any of the active ingredients.
- Do not use in foals under 2 month old and stallions.

PRECAUTIONS

Equimic® Plus has been formulated to be used exclusively in horses. It should not be used in other species.

SPECIAL PRECAUTIONS FOR USE

Equimic® Plus has been formulated to be used exclusively in horses. It should not be used in other species.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- In the event of accidental ingestion, do not induce vomiting. Consult to the doctor immediately.
- If contact with skin or eyes, wash with abundant water. If irritation occurs, consult to the doctor.
- Wash your hands after the administration of this product.

WARNINGS

Mantener fuera del alcance de los niños.

SIDE EFFECTS

ADVERSE EFFECTS

In the case of serious infestations, it can be observed signs of transient colic.

GUARD PERIOD

Do not administer to horses which meat is intended for human consumption.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:

Do not dispose product container, empty or with product along with the domestic waste.
Do not eliminate it on the ground or in water courses since the product is dangerous to aquatic organisms.

CONSERVATION

Store in a cool and dry place, between 15° and 30°C, protected from sunlight.
Once the syringe has been opened, use within 3 months.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

20 g graduated syringe.

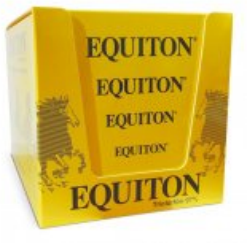
RECORDS

Chile: Reg. SAG No. 2171-B

EQUITON® ORAL POWDER.

POLVO ORAL.

ANTIPARASITARIO INTERNO Y EXTERNO.



Technical Specification

SPECIES

Horses.

It is recommended as internal anti-parasite for the treatment of gastrointestinal parasites in horses, in particular *Gasterophilus sp.* ("horse bot larvae" in the stomach).

DOSAGE FORM

Oral powder.

THERAPEUTIC ACTION

Internal and external anti-parasite.

COMPOSITION

Each 100 g of powder contains:
Trichlorfon .....100 g

INDICATIONS

- Do not administer in pregnant females (over 7 months of gestation) or in lactation.
- Do not administer in breeding animals.
- Do not administer in foals under 4 months.
- Do not administer in weakened animals.
- Do not administer to animals treated with other organophosphate compounds, anesthetics, muscle relaxants, until a period of 2 weeks after treatment.
- Do not administer to animals hypersensitive to organophosphates.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration way: Oral.

Dose:
Dose of the active principle: 40 mg of Trichlorfon / Kg of weight.
Product dose: 1 sachet of 20 grams for every 500 Kg of body weight, in a single dose.

How to use:

Dissolve the content of an EQUITON® sachet in 200 mL of water (1 cup) and administer orally.

DRUG INTERACTIONS

Trichlorfon should not be administered concomitantly with other organophosphates, with anesthetics and muscle relaxants.

CONTRAINDICATIONS

- Do not administer in pregnant females (over 7 months of gestation) or in lactation.
- Do not administer in breeding animals.
- Do not administer in foals under 4 months.
- Do not administer in weakened animals.
- Do not administer to animals treated with other organophosphate compounds, anesthetics, muscle relaxants, until a period of 2 weeks after treatment.
- Do not administer to animals hypersensitive to organophosphates.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not handle by pregnant or lactating women, or people who are hypersensitive to organophosphates.
- Avoid contact with food or fodder products.
- Potentially toxic product, if used negligently.
- Do not allow children to come into contact with treated animals.
- In humans, chronic exposure to organophosphates can cause damage to the nervous system and clinical signs such as headache, anxiety or irritability. It is recommended that the operators or handlers of this product use protective equipment (mask, gloves) to avoid inhalation or contact with the skin.

ANTIDOTE

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Do not administer to animals intended for human consumption.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
Dispose of this product with caution with household waste. Do not dispose of containers with product residues on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Box containing 25 packets of 20 g each.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG N° 945

EQUIVERM® ORAL PASTE

PASTA ORAL.

ANTIPARASITARIO

Technical Specification

SPECIES

Horses.

Infections from *Gasterophilus sp.* larvae,
Infections from worms: *Parascaris equorum, Oxyuris equi, Strongylus edentatus, Strongylus vulgaris, Strongylus equinus, Cyathostoma spp., Triodontophorus sp., Trichostrongylus axei, Oesophagodontus sp., Poteriostomun sp., Gyalocephalus capitatus*

DOSAGE FORM

Oral paste.

THERAPEUTIC ACTION

Antiparasitic.

COMPOSITION

Each gram contains:
Mebendazole.....100 mg
Trichlorfon.....459 mg
Excipients q.s.p.....1 g

PROPERTIES

Equiverm® is an effective product again *Gasterophilus* larvae and gastrointestinal nematodes in horses.

INDICATIONS

- Do not administer in fasting or empty stomach animals.
- This product should not be used simultaneously with other cholinesterase inhibitor anti-parasites, of internal or external use.
- As the product contains Trichlorfon, It is not recommended the administration to pregnant mares.

MODE OF APPLICATION

Place the product on the posterior part of the tongue. It is suitable for the horse, during the treatment, not to have anything that can be mixed with the product (for example, hay or other food).

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration.

Active ingredient dose:
8 mg of Mebendazole + 40 mg of Trichlorfon for each Kg of body weight. Single dose.

Product dose:
1 syringe of Equiverm® for 450 Kg of body weight. Single dose.

CONTRAINDICATIONS

- Do not administer in fasting or empty stomach animals.
- This product should not be used simultaneously with other cholinesterase inhibitor anti-parasites, of internal or external use.
- As the product contains Trichlorfon, It is not recommended the administration to pregnant mares.

WARNINGS

- Mantener fuera del alcance de los niños.

SIDE EFFECTS

The product is well tolerated at therapeutic doses. In some cases it can be observed mild cramps and soft stools. These side effects spontaneously disappear after a few hours. In the case of accidental overdose, it can cause typical toxic reactions of organophosphorus poisoning. Atropine can be used as an antidote for the symptoms. In case of poisoning, consult your nearest veterinarian.

GUARD PERIOD

Do not use the meat of treated animal for human consumption.

OBSERVATIONS

The consistence of the product depends on the temperature. It is necessary that the temperature be equal to or greater than 15ºC just before using it; in other case, the horse could reject the product due to excessive viscosity or by hardening that hinders its output. In very cold weather, it is recommended to warm the syringe before using it.

Nice apple flavor.

CONSERVATION

Store in a cool and dry place, at room temperature between 15º and 30ºC.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Syringe dispenser with 40 g of paste.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- **Chile:** Reg. SAG N° 069
- **Rep. Dominicana:** Reg. Nº 6092

EQUS® 20% - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANTIINFLAMATORIO - ANALGÉSICO.

Technical Specification

SPECIES

Horses.

Equs® is an anti-inflammatory and analgesic that can be used in musculoskeletal inflammations, mild claudications, soft tissue inflammations and trauma.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Anti-inflammatory - Analgesic.

COMPOSITION

Each 100 mL of solution contains:
Phenylbutazone.....20 g
Excipients q.s.p.....100 mL

INDICATIONS

- Do not administer to animals with heart, renal or hepatic impairment.
- Do not administer to pregnant or lactating females.
- Do not administer to dehydrated animals.
- Do not administer to animals with gastrointestinal mucous lesions.
- Do not administer to animals hypersensitive to the active ingredient.

ROUTE OD ADMINISTRATION AND DOSAGE

Intravenous administration.

Active ingredient dose:

- 2.2 – 4.4 mg/Kg of weight every 12 hours for 5 days.

Product dose:

- 0.5 –1 mL/45 Kg of weight every 12 hours for 5 days. The highest dose of 4.4 mg / Kg (1 mL / 45 Kg) should not be used beyond the first day of treatment.

DRUG INTERACTIONS

Anabolics inhibit the metabolism of Phenylbutazone.

CONTRAINDICATIONS

- Do not administer to animals with heart, renal or hepatic impairment.
- Do not administer to pregnant or lactating females.
- Do not administer to dehydrated animals.
- Do not administer to animals with gastrointestinal mucous lesions.
- Do not administer to animals hypersensitive to the active ingredient.

PRECAUTIONS

Special warnings and precautions for use:

- Keep out of the reach of children.
- In foals there is a greater risk of poisoning due to lack of maturity of the enzyme systems.
- Careful use is recommended in foals due to the higher incidence of hypoproteinemia and gastrointestinal ulceration.

SPECIAL PRECAUTIONS FOR USE

Special warnings and precautions for use:

- Keep out of the reach of children.
- In foals there is a greater risk of poisoning due to lack of maturity of the enzyme systems.
- Careful use is recommended in foals due to the higher incidence of hypoproteinemia and gastrointestinal ulceration.

SIDE EFFECTS

ADVERSE EFFECTS

The main reported adverse effects are gastrointestinal. Clinical signs include decreased appetite, depression, colic, weight loss, abdominal edema, hypoproteinemia, and diarrhea. Bleeding and ulcers can occur in the mouth, esophagus, stomach, intestine, cecum, and right dorsal colon.

GUARD PERIOD

Do not administer to horses which meat in intended for human consumption.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of the containers with the rest of the product on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 ° C, protected from light.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

100 mL vial

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG Nº 64
Rep. Dominicana: Reg. N° 11267

EQUS® ORAL TABLETS

COMPRIMIDO ORAL.

ANALGÉSICO, ANTIINFLAMATORIO NO ESTEROIDAL

Technical Specification

SPECIES

Horses.

Analgesic. Non-steroidal anti-inflammatory, indicated in the treatment of alterations in muscles, tendons, ligaments, joints and skin in horses.

DOSAGE FORM

Oral tablet.

THERAPEUTIC ACTION

Analgesic, Non-steroidal anti-inflammatory

COMPOSITION

Each Tablet contains:
Phenylbutazone.....1 g
Excipients q.s.p.....1 tablet

INDICATIONS

- Do not exceed a maximum dose of 4 g per horse per day.
- Do not administer in animals hypersensitive to phenylbutazone.
- Do not administer to animals with heart, kidney or liver failure, dehydrated animals, or in animals with hematological or bone marrow abnormalities.
- Do not administer in pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route of administration.

Product dose:

2 tablets for each 500 k of weight, every 12 hours the first day of treatment, followed by one tablet for every 500 Kg of weight, every 12 hours for 5 to 7 days.

DRUG INTERACTIONS

It is not recommended to use concomitantly with other pharmaceutical products.

CONTRAINDICATIONS

- Do not exceed a maximum dose of 4 g per horse per day.
- Do not administer in animals hypersensitive to phenylbutazone.
- Do not administer to animals with heart, kidney or liver failure, dehydrated animals, or in animals with hematological or bone marrow abnormalities.
- Do not administer in pregnant or lactating females.

PRECAUTIONS

- Do not exceed the established dose, since the therapeutic index of Phenylbutazone is low.
- Avoid as much as possible its use in foals or older animals due to the risk of kidney toxicity.

SPECIAL PRECAUTIONS FOR USE

- Do not exceed the established dose, since the therapeutic index of Phenylbutazone is low.
- Avoid as much as possible its use in foals or older animals due to the risk of kidney toxicity.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Wash your hands after handling the product. In case of accidental ingestion, do not induce vomiting. Get medical help.

WARNINGS

Mantener fuera del alcance de los niños.

SIDE EFFECTS

ADVERSE EFFECTS

Unwanted effects and adverse reactions:
Gastrointestinal adverse effects are the most important in phenylbutazone therapy in horses. Possible clinical signs include decreased appetite, depression, colic, weight loss, abdominal edema, hypoproteinemia (decreased blood protein), and diarrhea. In addition, bleeding and ulcers can occur in the mouth, esophagus, stomach, intestine, cecum, and right dorsal colon.

GUARD PERIOD

Do not administer to animals intended for human consumption.

OBSERVATIONS

Special precautions for disposal of waste material:
Discard any unused product remains in its original container. Dispose of the waste of this product with care together with household waste. Do not dispose of empty containers or the rest of the product on the ground or water courses.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° and 30°C.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

10 tablets

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG Nº: 1464
Perú: Reg. SENASA F.99.21.I.0114

COUNTRIES WHERE IT IS MARKETED

Imported and Distributed in Peru by:
Representaciones Durand SAC.
Av. Manuel Olguín N ° 501 Office N ° 604 Santiago de Surco Lima

EQUS® PLUS CREAM

CREMA.

ANTIINFLAMATORIO ANALGÉSICO.

Technical Specification

SPECIES

Horses.

It is recommended for the anti-inflammatory and analgesic treatment at the level of skin, subcutaneous tissue, muscle and joints in horses.

DOSAGE FORM

Cream.

THERAPEUTIC ACTION

Anti-inflammatory and analgesic.

COMPOSITION

Each 100 g of cream contains:
Phenylbutazone….....5.0 g
Lidocaine.…....2.0 g
Dimethyl sulfoxide.....10.0 g
Excipients q.s.p.....100 g

INDICATIONS

- Do not administer to animals with heart, kidney or liver failure, dehydrated, or with hematological or bone marrow abnormalities or that are hypersensitive to the active.
- Do not administer to pregnant or lactating females.

MODE OF APPLICATION

ROUTE OD ADMINISTRATION AND DOSAGE

Topical administration route.

Mode of use:
Clean the skin in the area to be applied, then apply a sufficient amount of product that allows to leave a thin layer on the affected area, apply two or three times a day massaging gently. Use for a maximum of 14 days or as indicated by the Veterinarian.

CONTRAINDICATIONS

- Do not administer to animals with heart, kidney or liver failure, dehydrated, or with hematological or bone marrow abnormalities or that are hypersensitive to the active.
- Do not administer to pregnant or lactating females.

PRECAUTIONS

Special warnings and precautions for use:

- Keep out of the reach of children.
- Dimethyl sulfoxide can cause irritation, erythema, vesicles, dry skin, or a local allergic reaction, which are transient and resolve rapidly when therapy is discontinued.
- Both Phenylbutazone and Dimethylsulfoxide can mask an existing pathology, due to their anti-inflammatory and analgesic properties.
- Use with caution in dehydrated animals or in animals in shock, since the diuretic and vasodilator effect of dimethylsulfoxide can exacerbate these conditions.
- In competition horses, use with caution due to the content of Lidocaine in its formulation.

SPECIAL PRECAUTIONS FOR USE

Special warnings and precautions for use:

- Keep out of the reach of children.
- Dimethyl sulfoxide can cause irritation, erythema, vesicles, dry skin, or a local allergic reaction, which are transient and resolve rapidly when therapy is discontinued.
- Both Phenylbutazone and Dimethylsulfoxide can mask an existing pathology, due to their anti-inflammatory and analgesic properties.
- Use with caution in dehydrated animals or in animals in shock, since the diuretic and vasodilator effect of dimethylsulfoxide can exacerbate these conditions.
- In competition horses, use with caution due to the content of Lidocaine in its formulation.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- When applying the product use protective gloves.
- If there is contact with skin, wash immediately with plenty of soap and water. In case of contact with the eyes, it is recommended to wash immediately with plenty of water for 15 minutes and in accidental ingestion rinse the mouth with plenty of water. Do not induce vomiting unless directed by a doctor. It is recommended to go to a healthcare center promptly.

WARNINGS

GUARD PERIOD

Do not use in animals intended for human consumption.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° and 30°C and protected from sunlight.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Knob with 120 g

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG Nº 0375
- Rep. Dominicana: Reg. Nº 5599
- Panamá: Reg. Nº RF-4130-18

FATROXIMIN® INTRAMAMMARY SUSPENSION

UNGÜENTO INTRAMAMARIO.

ANTIMASTÍTICO PARA EL TRATAMIENTO EN SECADO.

Technical Specification

SPECIES

Bovines (cows in dry period).

FATROXIMIN®, intramammary ointment, is indicated in the drying of the cow for the therapy of existing subclinical mastitis, the prevention of a possible infection during the dry period and the prevention of acute postpartum mastitis, produced by Gram positive microorganisms (*Streptococcus spp. , Staphylococcus spp., Corynebacterium spp.*) And Gram negative (*E. coli*).
It acts on the most frequent microbial species in the udder sensitive to Rifaximin, such as: *Streptococcus agalatiae, Streptococcus dysgalactiae, Streptococcus uberis, Streptococcus faecalis, Staphylococcus aureus* (including the penicillin-resistant strains) and *Staphymidis epidermis.*

DOSAGE FORM

Intramammary ointment.

THERAPEUTIC ACTION

Antimastitic for drying treatment

COMPOSITION

Each 5 mL syringe tube contains:
Rifaximin ..... 0.100 g
Excipients q.s.p ..... 5 mL

PROPERTIES

FATROXIMIN®, intramammary ointment is a preparation based on Rifaximin, a new antibiotic obtained by original synthesis, belonging to the Rifamycin family.
The particular chemical structure has given the molecule such chemical-physical characteristics that it has a pharmacokinetic that is completely different from the other rifamycins currently on the market.
The studies carried out reveal a practically null passage through the intramammary epithelium, allowing optimal availability of Rifaximin at the level of the fourth treated.

FATROXIMIN®, intramammary ointment has a high antibacterial activity of the bactericidal type against Gram positive microorganisms (Streptococcus spp., Staphylococcus spp., Corynebacterium spp.) And Gram negative ones (E. coli).
It acts on the most frequent microbial species in the udder.

INDICATIONS

- Do not administer in animals with hypersensitivity to Rifaximin.
- Do not administer in lactating cows.
- Do not administer to cows with clinical mastitis.

EFFECTIVENESS

CLINICAL EFFECTIVENESS
Field tests have shown:

- A high therapeutic activity, with bacteriological negativization of the rooms subjected to treatment and previously found infected due to *Streptococcus agalatiae, Streptococcus dysgalactiae, Streptococcus faecalis, Staphylococcus aureus, Staphylococcus epidermidis*, with some penicillin resistant strains.
- A consequent preventive action, especially in secretory disorders of the breasts, that is, in those pathophysiological situations characterized by a high cellular content with the absence of pathological microorganisms in the milk.

ROUTE OD ADMINISTRATION AND DOSAGE

Intramammary route of administration

- Administer a 5 mL syringe tube of FATROXIMIN®, intramammary ointment, (equivalent to 100 mg of Rifaximin) per breast quarter, after the last milking, prior to drying.
- Milk the quarter thoroughly and after disinfecting the nipple orifice, then apply FATROXIMIN®, intramammary ointment, introducing the cannula and injecting the entire contents of the syringe.
- Subsequently, the nipple should be massaged from the bottom up to spread the product throughout the room.

CONTRAINDICATIONS

- Do not administer in animals with hypersensitivity to Rifaximin.
- Do not administer in lactating cows.
- Do not administer to cows with clinical mastitis.

PRECAUTIONS

SPECIAL WARNINGS AND PRECAUTIONS FOR USE

- Keep out of the reach of children.
- Antimastitic treatment through a partial insertion (a few millimeters in the nipple canal) of the cannula in the nipple canal reduces the probability of new breast infections, since it prevents the dilation of the sphincter, the destruction of the keratin layer and also , deposits antibiotic along the nipple canal.
- If there are injuries to the nipple, or if the animals are very nervous, the complete insertion of the cannula can facilitate the work.

SPECIAL PRECAUTIONS FOR USE

SPECIAL WARNINGS AND PRECAUTIONS FOR USE

- Keep out of the reach of children.
- Antimastitic treatment through a partial insertion (a few millimeters in the nipple canal) of the cannula in the nipple canal reduces the probability of new breast infections, since it prevents the dilation of the sphincter, the destruction of the keratin layer and also , deposits antibiotic along the nipple canal.
- If there are injuries to the nipple, or if the animals are very nervous, the complete insertion of the cannula can facilitate the work.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not handle by people who are hypersensitive to Rifaximin.
- Do not smoke, eat or drink during the handling and administration of the product.
- Avoid contact with skin, eyes, or mouth.
- Use of gloves when administering the product.
- Wash hands after administering the product.
- In case of accidental ingestion, immediately go to a medical center and show the product label.
- In case of contact with skin, eyes or mouth, wash immediately with plenty of running water.
- In case of skin irritation, go immediately to a medical center and show the product label.

GUARD PERIOD

Meat: 0 days.

Milk: 0 days.

Treatment must be done at least 42 days before scheduled delivery; under these conditions, no protection period is required. In the event of premature delivery, do not use the milk from 18 consecutive milkings for human consumption.

OBSERVATIONS

SPECIAL PRECAUTIONS FOR DISPOSAL OF UNUSED PRODUCT OR WASTE MATERIAL

- Empty containers can be discarded as household waste, without any special precautions.
- Do not dispose of containers with product residues on the ground or water courses.
- Contact the importing company or companies specialized in waste disposal, to receive recommendations on the disposal of expired or unused products.

CONSERVATION

Store the product between 2º and 30ºC, in a cool and dry place, protected from light.

PRODUCT ORIGIN

Italy

CONDITION OF SALE

Sale with Veterinary Medical prescription.

PRESENTATION

Case with 12 syringe tubes of 5 mL each.

PREPARED BY

Fatro S.p.A.- Italia.

RECORDS

Reg. SAG Nº 2075-B

FEBRECTAL® BOVINE - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANTIINFLAMATORIO ANALGÉSICO Y ANTIPIRÉTICO.

Technical Specification

SPECIES

Horses, cattle and pigs.

Anti-inflammatory, analgesic and antipyretic. Recommended to relieve inflammation, fever and pain associated with musculoskeletal disorders and relieve pain associated with colic.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Anti-inflammatory, analgesic and antipyretic.

COMPOSITION

Each mL of solution contains:
Flunixin Meglumine.....83 mg
(Equivalent to 50 mg of Flunixin)
Excipients q.s.p.....1 mL

INDICATIONS

- Do not use in animals with hypersensitivity to the active ingredient or other NSAIDs.
- Do not use in animals suffering from heart, liver or kidney disease.
- Do not use in animals with lesions of the gastrointestinal tract, such as ulcers and hemorrhages.
- Do not use when there are signs of blood dyscrasias or impaired hemostasis.
- Do not use in cases of colic caused by ileus and associated with dehydration.
- Do not use in animals suffering from chronic musculoskeletal disorders.
- Do not administer to animals under 72 hours of life.
- Do not administer to pregnant or lactating females or to breeding animals.

ROUTE OD ADMINISTRATION AND DOSAGE

- Horses: 1.1 mg/Kg/day (1 mL/45 Kg of weight) EV or IM, repeat for a maximum of 5 days.
- Cattle: 2.2 mg/Kg/day (2 mL/45 Kg of weight) IV, in one or two administrations.
- Pigs: 2.2 mg/Kg/day (2 mL/45 Kg of weight) IM for a maximum of 3 days.

DRUG INTERACTIONS

- Concurrent administration of potentially nephrotoxic drugs should be avoided.
- Flunixin can reduce the renal excretion of some drugs, increasing their toxicity.
- The drug should not be administered together with other NSAIDs or glucocorticoids, as it may increase the toxicity of both, especially at the gastrointestinal level, increasing the risk to suffer from gastrointestinal ulcers.

CONTRAINDICATIONS

- Do not use in animals with hypersensitivity to the active ingredient or other NSAIDs.
- Do not use in animals suffering from heart, liver or kidney disease.
- Do not use in animals with lesions of the gastrointestinal tract, such as ulcers and hemorrhages.
- Do not use when there are signs of blood dyscrasias or impaired hemostasis.
- Do not use in cases of colic caused by ileus and associated with dehydration.
- Do not use in animals suffering from chronic musculoskeletal disorders.
- Do not administer to animals under 72 hours of life.
- Do not administer to pregnant or lactating females or to breeding animals.

PRECAUTIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be discarded as domestic waste, without any special precautions. Nope
Dispose of containers with product residues on the ground or in watercourses. For products expired or not used contact the manufacturing laboratory.

SPECIAL PRECAUTIONS FOR USE

Special precautions for disposal of unused product or waste material:
Empty containers can be discarded as domestic waste, without any special precautions. Nope
Dispose of containers with product residues on the ground or in watercourses. For products expired or not used contact the manufacturing laboratory.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not eat, drink or smoke while handling the product.
- Avoid self-injection.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

- In horses after IM injection, cases of localized inflammation have been reported, induration, stiffness and sweating.
- Accidental intra-arterial injection may cause stimulation of the central nervous system with signs such as ataxia, hyperventilation and muscle weakness. Clinical signs are transient and They generally do not require any treatment.
- Flunixin is a relatively safe agent for use in the horse, but there is the possibility of Gastrointestinal intolerance, hypoproteinemia, and hematologic abnormalities may occur.
- In horses and cattle, unusual anaphylactic reactions have been reported, mainly after rapid IV administration.
- Hematochezia and hematuria have been reported in cattle treated longer than 3 day recommendation.
- In pigs, no adverse reactions have been reported, except for some cases of irritation at the point of injection.

GUARD PERIOD

- Do not use in horses intended for human consumption.
- Cattle: 7 days meat, 2 days milk.
- Pigs: 21 days meat.

CONSERVATION

Store between 15° and 30°C, protected from light. Once the container is opened, use within 28 days.
Discard unused product after that time period.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

50 mL vial

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 393
- Dominican Republic: Reg. No. 5602

FEBRECTAL® EQUINE - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANTIINFLAMATORIO, ANALGÉSICO Y ANTIPIRÉTICO.

Technical Specification

SPECIES

Horses, cattle and pigs.

Anti-inflammatory, analgesic and antipyretic. Recommended to relieve inflammation, fever and pain associated with musculoskeletal disorders and relieve pain associated with colic.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Anti-inflammatory, analgesic and antipyretic.

COMPOSITION

Each mL of solution contains:

Flunixin Meglumine.....83 mg

(Equivalent to 50 mg of Flunixin)

Excipients q.s.p.....1 mL

INDICATIONS

- Do not use in animals with hypersensitivity to the active ingredient or other NSAIDs.
- Do not use in animals suffering from heart, liver or kidney disease.
- Do not use in animals with lesions of the gastrointestinal tract, such as ulcers and hemorrhages.
- Do not use when there are signs of blood dyscrasias or impaired hemostasis.
- Do not use in cases of colic caused by ileus and associated with dehydration.
- Do not use in animals suffering from chronic musculoskeletal disorders.
- Do not administer to animals under 72 hours of life.
- Do not administer to pregnant or lactating females or to breeding animals.

ROUTE OD ADMINISTRATION AND DOSAGE

- Horses: 1.1 mg/Kg/day (1 mL/45 Kg of weight) EV or IM, repeat for a maximum of 5 days.
- Cattle: 2.2 mg/Kg/day (2 mL/45 Kg of weight) IV, in one or two administrations.
- Pigs: 2.2 mg/Kg/day (2 mL/45 Kg of weight) IM for a maximum of 3 days.

DRUG INTERACTIONS

- Concurrent administration of potentially nephrotoxic drugs should be avoided. - Flunixin can reduce the renal excretion of some drugs, increasing their toxicity. - The drug should not be administered together with other NSAIDs or glucocorticoids, as it can increase the toxicity of both, especially at the gastrointestinal level, increasing the risk of suffering from gastrointestinal ulcers.

CONTRAINDICATIONS

- Do not use in animals with hypersensitivity to the active ingredient or other NSAIDs.
- Do not use in animals suffering from heart, liver or kidney disease.
- Do not use in animals with lesions of the gastrointestinal tract, such as ulcers and hemorrhages.
- Do not use when there are signs of blood dyscrasias or impaired hemostasis.
- Do not use in cases of colic caused by ileus and associated with dehydration.
- Do not use in animals suffering from chronic musculoskeletal disorders.
- Do not administer to animals under 72 hours of life.
- Do not administer to pregnant or lactating females or to breeding animals.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not eat, drink or smoke while handling the product.
- Avoid self-injection.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

- In horses after IM injection, cases of localized inflammation have been reported, induration, stiffness and sweating.
- Accidental intra-arterial injection may cause stimulation of the central nervous system with signs such as ataxia, hyperventilation and muscle weakness. Clinical signs are transient and They generally do not require any treatment.
- Flunixin is a relatively safe agent for use in the horse, but there is the possibility of Gastrointestinal intolerance, hypoproteinemia, and hematologic abnormalities may occur.
- In horses and cattle, unusual anaphylactic reactions have been reported, mainly after rapid IV administration.
- Hematochezia and hematuria have been reported in cattle treated longer than 3 day recommendation.
- In pigs, no adverse reactions have been reported, except for some cases of irritation at the point of injection.

GUARD PERIOD

- Do not use in horses intended to human consumption.
- Bovine: 7 days for meat, 2 days for milk.
- Pigs: 21 days for meat.

CONSERVATION

Store between 15° and 30°C, protected from light. Once the container is opened, use within 28 days. Discard unused product after that time period.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

20 and 50 mL ampule.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 393
- Dominican Republic: Reg. No. 5602

FIPRODRAG® 0,5 ML - CATS

SOLUCIÓN TÓPICA.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

FIPRODRAG® is indicated for the treatment and control of fleas (*Ctenocephalides felis*) and mites (*Cheyletiella spp.*) infestation in cats. FIPRODRAG® is also recommended as adjuvant therapy in flea allergy dermatitis (FAD) in cats.

Efficacy: With a single application, FIPRODRAG® allows to control flea reinfestation in cats, during a period of at least 5 weeks. If parasite burden is high, a monthly application of product is advisable.

For its lipophilic properties, FIPRODRAG® resists baths carried out 48 hours after its application. Do not bathe your pet 48 hours before and after applying the product.

DOSAGE FORM

Topical solution.

THERAPEUTIC ACTION

Topical solution.

COMPOSITION

Each mL of solution contains:

Fipronil.....100 mg

Excipients q.s.p.....1 mL

PROPERTIES

Effective action against fleas in cats.

INDICATIONS

- Do not use on kittens less than 12 weeks old.
- Do not use in rabbits.

MODE OF APPLICATION

1. Remove the upper marked tip of the pipette.
2. Put the end of the pipette directly on the animal skin, at scapulae (shoulders) level or at the base of neck, trying to separate the hair.
3. Press and pour the full content of pipette on 2 or 3 points, as indicated in the figure.

ROUTE OD ADMINISTRATION AND DOSAGE

For external use, through skin application.

Dose of active ingredient:

From 6.7 to 13.4 mg/kg body weight of the animal to be treated. In cats, apply one pipette of 0.5 mL.

CONTRAINDICATIONS

- Do not use on kittens less than 12 weeks old.
- Do not use in rabbits.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Precautions to be taken by the person administering the product:

- Avoid touching the pipette content directly with your fingers.
- Wash your hands with soap and water after applying and handling the product.
- Do not eat food, smoke, or drink beverages during the application of product.
- Avoid handling your pet immediately after the treatment, until the coat is completely dry.
- Discard in a safe way the empty pipettes. Do not dispose the empty pipettes into natural waterways.

WARNINGS

- Mantener fuera del alcance de los niños.
- Solamente para uso externo en gatos.
- El producto puede ser dañino si es ingerido.
- Producto tóxico para peces, aves y abejas.
- Producto inflamable.

SIDE EFFECTS

If the animal licks itself, a short period of hypersalivation may occur. If there is hypersensitivity or local irritation, it is recommended to not repeat the treatment.

OBSERVATIONS

- POISON.
- Flammable.

CONSERVATION

- Store in the original pack, at room temperature between 15 and 30°C, and in a fresh place.
- Keep out of heat sources.
- Do not store with food products.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

Box or Thermo-contracting Display containing 1 pipette of 0.5 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

SAG Reg. No. 2162

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

FIPRODRAG® 0,67 ML - DOGS UNTIL 10 KG OF BODY WEIGHT

SOLUCIÓN TÓPICA.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Dogs until 10 kg of body weight.

FIPRODRAG® is indicated for the treatment and control of fleas (*Ctenocephalides felis* and *Ctenocephalides canis*), lice (*Trichodectes canis*) and ticks (*Rhipicephalus sanguineus*) infestation in dogs over 10 weeks old. It is also recommended as adjuvant therapy in flea allergy dermatitis (FAD) in dogs. In this case, it is recommended a monthly application of product, including the animals cohabiting with them.

Efficacy: With a single application, FIPRODRAG® allows to control reinfestation in dogs, during a period of at least 8 weeks against fleas, and 5 weeks against ticks. If parasite burden is high, a monthly application of product is advisable. For its lipophilic properties, FIPRODRAG® resists baths carried out 48 hours after its application. Do not bathe your pet 48 hours before and after applying the product.

DOSAGE FORM

Topical solution.

THERAPEUTIC ACTION

External antiparasitic.

COMPOSITION

Each mL of solution contains:

Fipronil.....100 mg

Excipients q.s.p.....1 mL

PROPERTIES

Effective action against fleas, lice and ticks in dogs.

INDICATIONS

- Do not use on puppies less than 10 weeks old.
- Do not administer in rabbits.

MODE OF APPLICATION

1. Remove the upper marked tip of the pipette.
2. Put the end of the pipette directly on the animal skin, at scapulae (shoulders) level or at the base of neck, trying to separate the hair.
3. Press and pour the full content of pipette on 2 or 3 points, as indicated in the figure.

ROUTE OD ADMINISTRATION AND DOSAGE

For external use, through skin application.

The dose of Fipronil formulated as a topical solution for dogs and cats has been established between 6.7 and 13.4 mg/kg body weight of the animal to be treated. In dogs until 10 kg of body weight, apply a pipette of 0.67 mL

CONTRAINDICATIONS

- Do not use on puppies less than 10 weeks old.
- Do not administer in rabbits.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Precautions to be taken by the person administering the product:

- Avoid touching the pipette content directly with your fingers.
- Wash your hands with soap and water after applying and handling the product.
- Do not eat food, smoke, or drink beverages during the application of product.
- Avoid handling your pet immediately after the treatment, until the coat is completely dry.
- Discard in a safe way the empty pipettes. Do not dispose the empty pipettes into natural waterways.

WARNINGS

- Mantener fuera del alcance de los niños.
- Solamente para uso externo.
- El producto puede ser dañino si es ingerido.
- Producto tóxico para peces, aves y abejas.
- Producto inflamable.

SIDE EFFECTS

If the animal licks itself, a short period of hypersalivation may occur. If there is hypersensitivity or local irritation, it is recommended to not repeat the treatment.

OBSERVATIONS

- POISON.
- Flammable.

CONSERVATION

- Store in the original pack, at room temperature between 15 and 30°C, and in a fresh place.
- Keep out of heat sources.
- Do not store with food products.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

Box or Thermo-contracting Display containing 1 pipette of 0.67 mL for dogs up to 10 kg of body weight.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

SAG Reg. No. 2162

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

FIPRODRAG® 1,34 ML - DOGS BETWEEN 10 KG AND 20 KG OF BODY WEIGHT

SOLUCIÓN TÓPICA.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Dogs between 10 kg and 20 kg of body weight.

FIPRODRAG® is indicated for the treatment and control of fleas (*Ctenocephalides felis* and *Ctenocephalides canis*), lice (*Trichodectes canis*) and ticks (*Rhipicephalus sanguineus)* infestation in dogs over 10 weeks old. It is also recommended as adjuvant therapy in flea allergy dermatitis (FAD) in dogs. In this case, it is recommended a monthly application of product, including the animals cohabiting with them.

Efficacy: With a single application, FIPRODRAG® allows to control reinfestation in dogs, during a period of at least 8 weeks against fleas, and 5 weeks against ticks. If parasite burden is high, a monthly application of product is advisable. For its lipophilic properties, FIPRODRAG® resists baths carried out 48 hours after its application. Do not bathe your pet 48 hours before and after applying the product.

DOSAGE FORM

Topical solution.

THERAPEUTIC ACTION

External antiparasitic.

COMPOSITION

Each mL of solution contains:

Fipronil.....100 mg

Excipients q.s.p.....1 mL

PROPERTIES

Effective action against fleas, lice and ticks in dogs.

INDICATIONS

- Do not use on puppies less than 10 weeks old.
- Do not administer in rabbits.

MODE OF APPLICATION

- Remove the upper marked tip of the pipette.
- Put the end of the pipette directly on the animal skin, at scapulae (shoulders) level or at the base of neck, trying to separate the hair.
- Press and pour the full content of pipette on 2 or 3 points, as indicated in the figure.

ROUTE OD ADMINISTRATION AND DOSAGE

For external use, through skin application.

The dose of Fipronil formulated as a topical solution for dogs and cats has been established between 6.7 and 13.4 mg/kg body weight of the animal to be treated. In dogs between 10 kg and 20 kg of body weight, apply a pipette of 1.34 mL

CONTRAINDICATIONS

- Do not use on puppies less than 10 weeks old.
- Do not administer in rabbits.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Precautions to be taken by the person administering the product:

- Avoid touching the pipette content directly with your fingers.
- Wash your hands with soap and water after applying and handling the product.
- Do not eat food, smoke, or drink beverages during the application of product.
- Avoid handling your pet immediately after the treatment, until the coat is completely dry.
- Discard in a safe way the empty pipettes. Do not dispose the empty pipettes into natural waterways.

WARNINGS

- Mantener fuera del alcance de los niños.
- Solamente para uso externo.
- El producto puede ser dañino si es ingerido.
- Producto tóxico para peces, aves y abejas.
- Producto inflamable.

SIDE EFFECTS

If the animal licks itself, a short period of hypersalivation may occur.

If there is hypersensitivity or local irritation, it is recommended to not repeat the treatment.

OBSERVATIONS

- POISON.
- Flammable.

CONSERVATION

- Store in the original pack, at room temperature between 15 and 30°C, and in a fresh place.
- Keep out of heat sources.
- Do not store with food products.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

Box or Thermo-contracting Display containing 1 pipette of 1.34 mL for dogs from 10 to 20 kg of body weight.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

SAG Reg. No. 2162

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

FIPRODRAG® 2,68 ML - DOGS BETWEEN 20 KG AND 40 KG OF BODY WEIGHT

SOLUCIÓN TÓPICA.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Dogs between 20 kg and 40 kg of body weight.

FIPRODRAG® is indicated for the treatment and control of fleas (*Ctenocephalides felis* and *Ctenocephalides canis*), lice (*Trichodectes canis*) and ticks (*Rhipicephalus sanguineus*) infestation in dogs over 10 weeks old. It is also recommended as adjuvant therapy in flea allergy dermatitis (FAD) in dogs. In this case, it is recommended a monthly application of product, including the animals cohabiting with them.

Efficacy: With a single application, FIPRODRAG® allows to control reinfestation in dogs, during a period of at least 8 weeks against fleas, and 5 weeks against ticks. If parasite burden is high, a monthly application of product is advisable. For its lipophilic properties, FIPRODRAG® resists baths carried out 48 hours after its application. Do not bathe your pet 48 hours before and after applying the product.

DOSAGE FORM

Topical solution.

THERAPEUTIC ACTION

External antiparasitic.

COMPOSITION

Each mL of solution contains:

Fipronil.....100 mg

Excipients q.s.p.....1 mL

PROPERTIES

Effective action against fleas, lice and ticks in dogs.

INDICATIONS

- Do not use on puppies less than 10 weeks old.
- Do not administer in rabbits.

MODE OF APPLICATION

1. Remove the upper marked tip of the pipette.
2. Put the end of the pipette directly on the animal skin, at scapulae (shoulders) level or at the base of neck, trying to separate the hair.
3. Press and pour the full content of pipette on 2 or 3 points, as indicated in the figure.

ROUTE OD ADMINISTRATION AND DOSAGE

For external use, through skin application.

The dose of Fipronil formulated as a topical solution for dogs and cats has been established between 6.7 and 13.4 mg/kg body weight of the animal to be treated. In dogs between 20 kg and 40 kg of body weight, apply a pipette of 2.68 mL

CONTRAINDICATIONS

- Do not use on puppies less than 10 weeks old.
- Do not administer in rabbits.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Precautions to be taken by the person administering the product:

- Avoid touching the pipette content directly with your fingers.
- Wash your hands with soap and water after applying and handling the product.
- Do not eat food, smoke, or drink beverages during the application of product.
- Avoid handling your pet immediately after the treatment, until the coat is completely dry.
- Discard in a safe way the empty pipettes. Do not dispose the empty pipettes into natural waterways.

WARNINGS

- Mantener fuera del alcance de los niños.
- Solamente para uso externo.
- El producto puede ser dañino si es ingerido.
- Producto tóxico para peces, aves y abejas.
- Producto inflamable.

SIDE EFFECTS

- If the animal licks itself, a short period of hypersalivation may occur.
- If there is hypersensitivity or local irritation, it is recommended to not repeat the treatment.

OBSERVATIONS

- POISON.
- Flammable.

CONSERVATION

- Store in the original pack, at room temperature between 15 and 30°C, and in a fresh place.
- Keep out of heat sources.
- Do not store with food products.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

Box or Thermo-contracting Display containing 1 pipette of 2.68 mL for dogs from 20 to 40 Kg of body weight.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

SAG Reg. No. 2162

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

FIPRODRAG® 4,02 ML - DOGS BETWEEN 40 KG AND 60 KG OF BODY WEIGHT

SOLUCIÓN TÓPICA.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Dogs between 40 kg and 60 kg of body weight.

FIPRODRAG® is indicated for the treatment and control of fleas (*Ctenocephalides felis* and *Ctenocephalides canis*), lice (*Trichodectes canis*) and ticks (*Rhipicephalus sanguineus*) infestation in dogs over 10 weeks old. It is also recommended as adjuvant therapy in flea allergy dermatitis (FAD) in dogs. In this case, it is recommended a monthly application of product, including the animals cohabiting with them.

Efficacy: With a single application, FIPRODRAG® allows to control reinfestation in dogs, during a period of at least 8 weeks against fleas, and 5 weeks against ticks. If parasite burden is high, a monthly application of product is advisable. For its lipophilic properties, FIPRODRAG® resists baths carried out 48 hours after its application. Do not bathe your pet 48 hours before and after applying the product.

DOSAGE FORM

Topical solution.

THERAPEUTIC ACTION

External antiparasitic.

COMPOSITION

Each mL of solution contains:

Fipronil.....100 mg

Excipients q.s.p.....1 mL

PROPERTIES

Effective action against fleas, lice and ticks in dogs.

INDICATIONS

- Do not use on puppies less than 10 weeks old.
- Do not administer in rabbits.

MODE OF APPLICATION

1. Remove the upper marked tip of the pipette.
2. Put the end of the pipette directly on the animal skin, at scapulae (shoulders) level or at the base of neck, trying to separate the hair.
3. Press and pour the full content of pipette on 2 or 3 points, as indicated in the figure.

ROUTE OD ADMINISTRATION AND DOSAGE

For external use, through skin application.

The dose of Fipronil formulated as a topical solution for dogs and cats has been established between 6.7 and 13.4 mg/kg body weight of the animal to be treated. In dogs between 40 kg and 60 kg of body weight, apply a pipette of 4.02 mL

CONTRAINDICATIONS

- Do not use on puppies less than 10 weeks old.
- Do not administer in rabbits.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Precautions to be taken by the person administering the product:

- Avoid touching the pipette content directly with your fingers.
- Wash your hands with soap and water after applying and handling the product.
- Do not eat food, smoke, or drink beverages during the application of product.
- Avoid handling your pet immediately after the treatment, until the coat is completely dry.
- Discard in a safe way the empty pipettes. Do not dispose the empty pipettes into natural waterways.

WARNINGS

- Mantener fuera del alcance de los niños.
- Solamente para uso externo.
- El producto puede ser dañino si es ingerido.
- Producto tóxico para peces, aves y abejas.
- Producto inflamable.

SIDE EFFECTS

If the animal licks itself, a short period of hypersalivation may occur. If there is hypersensitivity or local irritation, it is recommended to not repeat the treatment.

OBSERVATIONS

- POISON.
- Flammable.

CONSERVATION

- Store in the original pack, at room temperature between 15 and 30°C, and in a fresh place.
- Keep out of heat sources.
- Do not store with food products.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

Box or Thermo-contracting Display containing 1 pipette of 4.02 mL for dogs from 40 to 60 kg of body weight.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

SAG Reg. No. 2162

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

FIPROKILL® SPRAY - SPRAY SOLUTION

SOLUCIÓN TÓPICA.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Dogs and cats.

FIPROKILL® SPRAY is effective in the control and treatment of flea (*Ctenocephalides felis*) and tick (*Rhipicephalus sanguineus*) infestation in dogs and cats from 2 days of age, providing a rapid knock-down effect against adult fleas.

DOSAGE FORM

Topical Solution

THERAPEUTIC ACTION

External antiparasitic.

COMPOSITION

Each 100 mL of product contains:
Fipronil.....0.25 g
Excipients q.s.p.....100 mL

PROPERTIES

INDICATIONS

- Do not apply to rabbits.
- Do not apply in animals with known hypersensitivity to fipronil.

EFFECTIVENESS

FIPROKILL® SPRAY in dogs provides a protection period of up to 3 months against fleas and up to 5 weeks against ticks. In cats, the period of protection against fleas extends for 4 weeks.

MODE OF APPLICATION

1. Apply the product in ventilated environments and make sure to use gloves. Avoid contact with skin. If there is contact with hands, wash with plenty of water after handling the product.
2. Apply always against the grain, as indicated in the figure, from a distance of approximately 20 cm.
3. The areas where the product should be applied are the belly and the flanks (chest, abdomen, armpits, shoulders and paws).
4. For long coat animals, separate the hair and apply the product “against the grain”, until the root.
5. Rub the coat for achieving the product penetrates the skin.
6. After applying, do not expose the pet to heat sources or fire. Let dry naturally. Do not use hairdryer.
7. For treating the face, pour a little of the product in the palm of the gloved hand, and then apply it rubbing gently the face, avoiding touching the eyes, mouth and mucous membranes.
8. Long or thick coat animals may require the application of a larger quantity of product.
9. It is recommended to not bathe the pet within 48 hours before and after applying Fiprokill® Spray.
10. The product is safe in kittens and puppies from 2 days of age, provided that the application is carried out following the manufacturer’s instructions.

ROUTE OD ADMINISTRATION AND DOSAGE

Dose of active ingredient:
7.5 mg - 15 mg/kg of body weight.

Dosage of product:

- **Fiprokill Spray 50 mL and 100 mL bottle:** Apply FIPROKILL® Spray Solution, approximately 6 to 12 sprays by kg of body weight (wetting the hair root), in such a way the animal becomes totally wet.
- **Fiprokill Spray 250 mL bottle:** Apply FIPROKILL® Spray Solution, approximately 2 to 4 sprays by kg of body weight (wetting the hair root), in such a way the animal becomes totally wet.

Long or thick coat animals may require the application of a larger quantity of product.

CONTRAINDICATIONS

- Do not apply to rabbits.
- Do not apply in animals with known hypersensitivity to fipronil.

WARNINGS

- Mantener fuera del alcance de los niños.
- Administrar con precaución en animales debilitados y seniles.
- Producto inflamable. Mantener alejado de las fuentes de calor

ADVERSE EFFECTS

- If the animal licks itself, there may be a brief period of hypersalivation (mainly in cats).
- Transient skin reactions at the application site such as erythema, itching, or alopecia may occur.
- In the event of hypersensitivity or local irritation, it is recommended not to repeat the treatment.

OBSERVATIONS

Frequently asked questions:

What happen if my pet licks itself after the application? What would happen to my pet?

FIPROKILL® SPRAY is formulated with an active ingredient that is safe for mammals; however, after applying the product, avoid your pet licks itself until its coat is dry.

Can I let my children caress the animals after the application? How long should I wait for?

It is advisable the pet is totally dry before touching it again. Otherwise, wash the hands with plenty of water.

Can I bathe my pet without altering the efficacy of FIPROKILL® SPRAY?

It is recommended to not bathe your pet at least 48 hours after applying the product.

CONSERVATION

Store between 15 and 30 °C, protected from light. Once the container is opened, use within 12 weeks. Discard unused product after that time period

CONDITION OF SALE

OTC product (non-prescription)

PRESENTATION

Bottle with spray valve with 50 mL, 100 mL and 250 mL

PREPARED BY

Drag Pharma Laboratory

RECORDS

Chile: Reg. SAG N° 1872
Panamá: Reg. Nº RF-4541-19
Rep. Dominicana: Reg. Nº 8664
Bolivia: Reg. SENASAG PUV-F N° 007256/16
Perú: Registro SENASA F.87.30.I.0193
Uruguay: Reg. MGAP N° 2020A00783

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

Imported and Distributed in Peru by:

Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

Importer in Uruguay: VIVAFIL S.A.

Av. 18 de Julio 1746 apt 601 - Montevideo - Uruguay.
grupotecnovet@gmail.com Technical Director: DMTV Diego Cuadrado.

NOTE

Special precautions for disposal of unused product or waste material: Dispose of this product with care together with household waste. Do not dispose of containers with product residues on the ground or in watercourses, as the product is toxic to birds, bees and fish. For expired or unused products, contact the manufacturing laboratory.

FIRE ENERGY®

SOLUCIÓN ORAL.

SUPLEMENTO NUTRICIONAL PARA CABALLOS.

Technical Specification

SPECIES

Horses.

THERAPEUTIC ACTION

Nutritional Suplement for horses.

INGREDIENTS

Glucose, Honey, Ginseng extract, Eucalyptus Oil, Menthol, Excipients.

TECHNICAL CHARACTERISTICS

Fire Energy® is a specially formulated product for high performance horses, providing an instant energy boost when needed.

The highly palatable combination of ingredients, which include Honey, Glucose and Ginseng as well as being an ideal energy source for horses in training or undergoing physical effort, stimulates the animal's appetite, making it an indispensable supplement for finicky horses.

USE INSTRUCTIONS

Administer 50 mL 2 -3 times a day per oral route.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Keep in a cool and dry place protected from light, below 30°C.

CONDITION OF SALE

OTC product (non-prescription).

RECORDS

Chile: Reg. SAG N°: RM 03-008N

NOTE

"VETERINARY USE ONLY”

"EXCLUSIVE USE IN ANIMAL FEEDING"

“THIS PRODUCT IS NOT A COMPLETE FEED"

FLOVOVERMIC® LARGE BREED - ORAL TABLET

COMPRIMIDO ORAL.

ANTIPARASITARIO INTERNO DE AMPLIO ESPECTRO.

Technical Specification

SPECIES

Large breed dogs (35 Kg).

Broad spectrum internal antiparasitic in a single dose. Effective against nematodes and cestodes, such as: *Toxocara canis, Toxascaris leonina, Trichuris vulpis, Ancylostoma caninum, Uncinaria stenocephala, Trichostrongilus sp., Taenia hydatigena, Taenia pisiformis, Taenia multiceps, Taenia serialis, Diphmis taenia serialis, and Diphmis taenia serialis canothinuminumidiforum, and Taenia serialis canothinumuminumidiforum. granulosus.*

DOSAGE FORM

Oral Tablet.

THERAPEUTIC ACTION

Broad spectrum internal anti-parasite.

COMPOSITION

Each tablet contains:
Flubendazole..... 770 mg
Praziquantel.....175 mg
Pyrantel Pamoato.....504 mg
(Equivalent to 175 mg Pyrantel)
Excipients q.s.p.....1 tablet

INDICATIONS

- Do not administer to dogs under 4 weeks of age.
- Do not administer to pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route, directly into the mouth or mixed with the food.

Dosage of active ingredients:
- Flubendazole: 22 mg / Kg of weight, in a single dose.
- Pyrantel: 5 mg / Kg of weight, in a single dose.
- Praziquantel: 5 mg / Kg of weight, in a single dose.

Product dosage:
1 tablet every 35 Kg of weight in a single dose. For the treatment of Trichuris vulpis infestations, administer 1 tablet every 24 hours, for 3 consecutive days.

CONTRAINDICATIONS

- Do not administer to dogs under 4 weeks of age.
- Do not administer to pregnant or lactating females.

WARNINGS

- Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

Adverse effects are rarely observed and if they occur they are mild and transitory, with gastrointestinal symptoms such as nausea, vomiting, anorexia, abdominal pain and diarrhea being observed.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of water. Dispose of the waste of this product with care together with household waste.

CONSERVATION

Storage in a cool and dry place, at room temperature between 15° y 30°C, protected from sunlight.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Box with 20 tablets.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG Nº: 1390

Perú: Registro SENASA F.08.21.I.1058

Bolivia: Reg. SENASAG PUV- Nº 10003/21

COUNTRIES WHERE IT IS MARKETED

Imported and Distributed in Peru by:

Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

FLOVOVERMIC® ORAL TABLETS

COMPRIMIDO ORAL.

ANTIPARASITARIO INTERNO DE AMPLIO ESPECTRO EN DOSIS ÚNICA.

Technical Specification

SPECIES

Dogs and cats.

Broad spectrum internal anti-parasite, single dose. Effective against nematodes and cestodes.

DOSAGE FORM

Oral tablet.

THERAPEUTIC ACTION

Broad spectrum internal anti-parasite, in single dose.

COMPOSITION

Each tablet contains:
Flubendazole..... 220 mg
Praziquantel….....50 mg
Pyrantel Pamoate.....144 mg
(Equivalent to 50 mg Pyrantel base)
Excipients q.s.p.....1 tablet.

INDICATIONS

Do not administer to dogs younger than 4 weeks, nor to cats younger than 6 weeks of age.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration orally, directly in the oral cavity or mixed in food.

Dose:
Dogs and cats: Administer 1 comp./10 kg of body weight, in a single dose (equivalent to 22 mg / kg of Flubendazol, 5 mg / kg of Pirantel and 5 mg / kg of Praziquantel).

In dogs over 40 kilos in weight, 4 tablets are enough.

CONTRAINDICATIONS

Do not administer to dogs younger than 4 weeks, nor to cats younger than 6 weeks of age.

PRECAUTIONS

Special warnings, interactions and precautions for use:

- Keep out of the reach of children.
- Do not use concomitantly with other pharmaceutical products.

SPECIAL PRECAUTIONS FOR USE

Special warnings, interactions and precautions for use:

- Keep out of the reach of children.
- Do not use concomitantly with other pharmaceutical products.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Wash hands after administering the product.
- If accidental ingestion of the tablet occurs, consult a doctor, presenting the package.
- Do not smoke, eat or drink during the administration of the product.

WARNINGS

ADVERSE EFFECTS

Unwanted effects and adverse reactions:
Mild and transient gastrointestinal symptoms, such as nausea, vomiting, anorexia, abdominal pain, and / or diarrhea, may occur in some animals.

OBSERVATIONS

Precautions for disposing of unused product or waste material:

- Discard the remains of unused product in its original container.
- Do not throw the empty container or with product remains, in rivers, lakes or streams of natural water.
- Dispose of this product with caution with household waste.

CONSERVATION

Store at room temperature between 15° and 30°C.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Box with 60 tablets.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: SAG Reg. No. 0503
Panama: Reg. No RF-4185-18
Bolivia: Reg. SENASAG PUV-F-N° 005516/13
Imported and distributed by:
ZOO PHARMA VETERINARY INPUTS S.R.L.
Diaz Romero N° 1339 Miraflores Zone, La Paz-Bolivia
Telephone: 591 2223357
Costa Rica: Reg. No. MV-7109

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

F-L-T® SHAMPOO

SHAMPOO.

SHAMPOO

Technical Specification

SPECIES

Dogs and cats.

Complementary treatment, indicated in dermatopathies where an antimicrobial, antipruritic, acaricidal and keratolytic effect is necessary; seborrhea, pyoderma, eczema, dermatitis, scabies, dermatomycosis and in periodic preventive hygiene.

DOSAGE FORM

Shampoo.

THERAPEUTIC ACTION

Shampoo.

COMPOSITION

Each 100 ml contains:
Benzoyl Peroxide.....2.5 g
Sulfur..... 2.0 g
Excipients q.s.p.....100 mL

INDICATIONS

- Do not use in animals that present pictures of dry seborrhea or severe scaling.
- Do not administer to animals with hypersensitivity to the active principles.
- Do not use in pregnant or lactating females.

MODE OF APPLICATION

- Bathe beforehand with a neutral shampoo.
- Apply F-L-T in sufficient quantity to achieve abundant lather.
- Rub 2-3 minutes.
- Let stand for 5 minutes and rinse thoroughly.

The frequency and duration of the treatment will depend on the treating Veterinarian.

CONTRAINDICATIONS

- Do not use in animals that present pictures of dry seborrhea or severe scaling.
- Do not administer to animals with hypersensitivity to the active principles.
- Do not use in pregnant or lactating females.

PRECAUTIONS

Special warnings and precautions for use

- Keep out of the reach of children.
- For external use only.
- Avoid contact of the product with the animal's eyes and mouth.
- Avoid licking your hair after treatment.
- If any degree of irritation develops, discontinue treatment and consult a Veterinarian.

SPECIAL PRECAUTIONS FOR USE

Special warnings and precautions for use

- Keep out of the reach of children.
- For external use only.
- Avoid contact of the product with the animal's eyes and mouth.
- Avoid licking your hair after treatment.
- If any degree of irritation develops, discontinue treatment and consult a Veterinarian.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Avoid contact with the eyes, if this occurs, rinse with plenty of water.
- The product may cause fabric bleaching or jewelry staining.

ADVERSE EFFECTS

- Some adverse reactions observed after the administration of benzoyl peroxide are hyperemia and blistering.
- Animals with sensitive skin may develop excessive dryness or irritation.
- The product could stain the animal coat clear.

OBSERVATIONS

Special precautions for disposal of unused product or waste material.
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 ° and 30 ° C, protected from light.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

150 mL bottle.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG N° 940
Costa Rica: Reg. Nº MAG CL4-45-9-5650
Bolivia: Reg. SENASAG PUV-F-Nº 006140/14

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

FURASEP® OINTMENT

POMADA.

ANTISÉPTICO.

Technical Specification

SPECIES

Horses.

Recommended as an antiseptic for wounds, burns, abrasions and skin sores.

DOSAGE FORM

Ointment.

THERAPEUTIC ACTION

Antiseptic.

COMPOSITION

Each 100 g contains:

Nitrofurazone.....0.2 g

Excipients q.s.p.....100 g

USE INSTRUCTIONS

Apply enough ointment directly on the injury or soak a piece of gauze with Furasep®. The Furasep® application can be repeated several times a day; in addition, the preparation should remain on the injury not less than 24 hours and until the wound is closed.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Do not use in animals intended for human consumption.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° and 30°C, protected from sunlight.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

500 g container.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No. 1473

FURODRAG® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

DIURÉTICO.

Technical Specification

SPECIES

Horses and bovines.

Diuretic, recommended for the treatment of acute pulmonary edema, mammary, pregnancy, congestive heart failure and ascites; as well as supportive therapy in forced diuresis in case of intoxication of different origin. It is also recommended in cases of renal failure with oliguria and procedures of diagnostics. In nose bleeding horses (epistaxis).

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Diuretic.

COMPOSITION

Each ml of injectable solution contains:

Furosemide.....50 mg

Excipients q.s.p.....1 mL

ROUTE OD ADMINISTRATION AND DOSAGE

Intravenous or intramuscular route.

Recommended dose:

Bovine and horses: 5-10 ml/animal, once or twice a day.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Bovine:

Meat and milk: 48 hours.

Do not administer to horses which meat is intended for human consumption.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° and 30°C, protected from sunlight.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Ampule with 50 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 1500
- Dominican Republic: Reg. No. 5598

GASTROENTERIL® ORAL SUSPENSION

SUSPENSIÓN ORAL.

ANTIMICROBIANO - ANTIPROTOZOARIO.

Technical Specification

SPECIES

Dogs and cats.

Gastroenteril® oral suspension, is indicated for the treatment of diarrhea associated with Giardias, in dogs and cats. It is also indicated in enteric and systemic infections caused by obligate anaerobic bacteria sensitive to Metronidazole.

DOSAGE FORM

Oral suspension.

THERAPEUTIC ACTION

Antimicrobial - Antiprotozoal.

COMPOSITION

Each 100 ml of the product contains:
Metronidazole.....2.5 g
Excipients q.s.p.....100 mL

INDICATIONS

- Do not administer to patients with hypersensitivity to Metronidazole or nitroimidazole derivatives.
- Do not use during pregnancy, lactation or in breeding animals.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route.

Giardiasis and anaerobic infections:

- Active ingredient dose: 25 mg of Metronidazole / Kg, every 12 hours for 5-7 days *.
- Product dose: 1 mL / Kg of weight, twice a day for 5 to 7 days *.

* In the case of systemic anaerobic infections, therapy can be prolonged for 6 weeks.

DRUG INTERACTIONS

- Metronidazole may prolong prothrombin time in patients taking Warfarin or other coumarin anticoagulants.
- Phenobarbital or Phenytoin may increase the metabolism of Metronidazole and increase the likelihood of dose related side effects.

CONTRAINDICATIONS

- Do not administer to patients with hypersensitivity to Metronidazole or nitroimidazole derivatives.
- Do not use during pregnancy, lactation or in breeding animals.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Wash hands after use.
- Do not handle by people hypersensitive to Metronidazole.
- In case of contact with skin, eyes or mucosa, immediately wash with plenty of water.
- Do not handle by pregnant women.
- In case of accidental ingestion, immediately go to a medical center and show the product label.
- Do not eat, smoke or drink during its administration.

WARNINGS

Advertencias y precauciones especiales de uso:

- Gastroenteril® suspensión oral debe usarse con precaución en animales con disfunción hepática. En caso de deterioro hepático significativo, utilizar el 25-50% de la dosis usual.
- Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

Vomiting, hepatotoxicity, neutropenia. In prolonged treatments, neurological signs, which completely subside once the use of the product has been discontinued.

OBSERVATIONS

Shake vigorously before use.

Special precautions for the disposal of unused product or waste material:
Empty containers can be discarded as household waste without any special precautions. Do not dispose of the containers with the rest of the product on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

- Store at a temperature between 2 and 30 ° C, protected from light.
- Once opened, use the product within 3 months. Discard the unused product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

120 ml bottle.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG N° 2036
Costa Rica: Reg. N° MAG CL4-42-18-5785
Uruguay: Reg. MGAP N° 2016A00378
Bolivia: Reg. SENASAG PUV-F-N° 006142/14
Perú: Registro SENASA F.91.02.I.0005

COUNTRIES WHERE IT IS MARKETED

Uruguay:
Importador:
VIVAFIL S.A.
RIO NEGRO 1107 Montevideo - Uruguay,
TEL 29001112
grupotecnovet@gmail.com
Director Técnico: DMTV Diego Cuadrado.

Bolivia:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

Perú:
Importado y Distribuido por Representaciones Durand SAC.
Av. Manuel Olguín N° 501 Oficina N° 604 Santiago de Surco Lima.

GLICEFAR® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

RELAJANTE MUSCULAR CENTRAL.

Technical Specification

SPECIES

Horses.

Adjuvant for the induction of anesthesia in horses.
It can be used as an inducing agent in combination with barbiturates to soften the stage of induction and recovery of the latter.
It can also be used in a combination with other agents such as Xylazine and Ketamine, to maintain anesthesia and thus decrease the concentration of inhalation anesthesia.
Its use is recommended only in adult horses.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Central muscle relaxant

COMPOSITION

Each mL of product contains:
Guaifenesin.....100 mg
Excipients q.s.p.....1 mL

INDICATIONS

Do not administer to pregnant or lactating females.

USE INSTRUCTIONS

- To use in combination with barbiturates, add the barbiturate (example: Thiopental) dose to the Glicefar® solution before the administration and proceed to the infusion or administer it as a bolus immediately after the administration of Glicefar®.
- To use in combination with xylazine and/or ketamine, Xylazine is administered 2 to 5 minutes before the administration of Glicefar® in the recommended dose and later the ketamine will be administered, once the ataxia signs from Glicefar® are present.

ROUTE OD ADMINISTRATION AND DOSAGE

Intravenous administration only.

Recommended dose: 100 - 110 mg / Kg
Recommended practical dose: 1 mL / Kg of weight.

CONTRAINDICATIONS

Do not administer to pregnant or lactating females.

PRECAUTIONS

- The extravascular injection should be avoided as it can cause irritation of the tissue.
- It is recommended to monitor the heart and breeding rate of the patient when administering Guaifenesin (Glyceryl Guaiacolate).

SPECIAL PRECAUTIONS FOR USE

- The extravascular injection should be avoided as it can cause irritation of the tissue.
- It is recommended to monitor the heart and breeding rate of the patient when administering Guaifenesin (Glyceryl Guaiacolate).

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Guaifenesin (Glyceryl Guaiacolate) is irritant to skin and mucous membranes, for that reason people should avoid any direct contact or accidental ingestion. The use of disposable gloves is recommended when handling this product.
- Wash hands after using the product.
- Guaifenesin (Glyceryl Guaiacolate) may cause hypersensitivity reactions.

ANTIDOTE

There is not specific antidote.

WARNINGS

Mantener fuera del alcance de los niños.

SIDE EFFECTS

ADVERSE EFFECTS

Symptoms of breeding apnea, nystagmus, hypotension and stiff muscles are associated with toxic drug levels.
One liter of the product can be administered without showing toxic effects in the individual.

GUARD PERIOD

Do not use in horses which meat is intended for human consumption.

CONSERVATION

Store in a cool and dry place, between 15° and 30°C, protected from sunlight.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

serum container with 500 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No. 2039
Costa Rica: Reg. N° MAG CL 4-3-11-6176
Perú: Reg. SENASA F.A0.01.I.0004

COUNTRIES WHERE IT IS MARKETED

Imported and Distributed by Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

HASYUN® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANTIINFLAMATORIO ESTEROIDAL.

Technical Specification

SPECIES

Cattle, pigs, horses, sheep, dogs and cats.

Recommended for the treatment of shock; inflammatory and allergic disorders in dogs, cats, cattle, horses and pigs; primary ketosis in cattle; delivery induction in cattle, sheep and pigs and Hypoadrenocorticism.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Steroidal anti-inflammatory.

COMPOSITION

Each ml of injectable solution contains:
Dexamethasone Sodium Phosphate.....5.26 mg
(Equivalent to 4 mg de Dexamethasone base)
Excipients q.s.p.....1 mL

INDICATIONS

- Do not use in viral infections.
- Do not use in animals with tuberculosis, chronic nephritis, Cushing syndrome, peptic ulcer or diabetes (unless used as emergency treatment).
- Do not use in animals hypersensitive to Dexamethasone.
- Do not administer to pregnant females.

ROUTE OD ADMINISTRATION AND DOSAGE

Intravenous and intramuscular route.

SPECIE	ACTIVE INGREDIENT DOSE ACCORDING SPECIE AND THERAPY	PRODUCT DOSE ACCORDING THERAPY
Dogs	Inflammatory disorders: 0.05-0.2 mg/Kg of weight a day, I.V. or I.M route, for 3-5 days. Shock: 4-6 mg/Kg of weight, I.V route, slowly, single dose.	Inflammatory disorders: 0.125-0.5 ml/10 Kg of weight a day, I.V. or I.M route, for 3-5 days. Shock: 1-1.5 ml/Kg of weight, I.V. route, slowly, single dose.
Cats	Inflammatory disorders: 0.1-0.3 mg/Kg of weight a day, I.V. or I.M route, for 3-5 days. Shock: 4-6 mg/Kg of weight, I.V route, slowly, single dose. ¡	Inflammatory disorders: 0.1-0.3 ml/4Kg of weight a day, I.V. or I.M route, for 3-5 days. Shock: 1-1.5 ml/Kg of weight, I.V. route, slowly, single dose.
Cattle	Inflammatory disorders, ketosis: 0.0125- 0.05 mg/Kg of weight a day, I.V. or I.M. route. Delivery induction: 0.05-0.075 mg/Kg of weight, I.M, route, single dose.	Inflammatory disorders, ketosis: 1.25-5 ml/400 Kg of weight a day, I.V. or I.M. route. Delivery induction: 5- 7.5 ml/400 Kg of weight, I.M. route, single dose.
Pigs	Inflammatory disorders: 0.02-0.08 mg/Kg of weight a day, I.V. or I.M route. Shock: 0.5-5 mg/Kg, I.V. route.	Inflammatory disorders: 0.5-2 ml/100 Kg of weight a day, I.V. or I.M. route. Shock: 0.125-1.25 mL/Kg, I.V. route.
Sheep	Delivery induction: 0.16-0.32 mg/Kg of weight, I.M, route, single dose.	Delivery induction: 2- 4 ml/50 Kg of weight, I.M., route, single dose.
Horses	Inflammatory disorders: 0.005-0.01 mg/Kg of weight a day, I.V. route. Shock: 4-6 mg/Kg, I.V. route.	Inflammatory disorders: 0.625-1.25 ml/500 Kg a day, I.V. route. Shock: 1 to 1.5 ml/Kg, I.V. route.

CONTRAINDICATIONS

- Do not use in viral infections.
- Do not use in animals with tuberculosis, chronic nephritis, Cushing syndrome, peptic ulcer or diabetes (unless used as emergency treatment).
- Do not use in animals hypersensitive to Dexamethasone.
- Do not administer to pregnant females.

PRECAUTIONS

Special warnings and precautions for use:

- Keep out of the reach of children.
- In case of prolonged therapy (more than 7 days), it should be gradually discontinued in successive days.
- If a permanent corticosteroid effect is required, oral therapy is recommended.

SPECIAL PRECAUTIONS FOR USE

Special warnings and precautions for use:

- Keep out of the reach of children.
- In case of prolonged therapy (more than 7 days), it should be gradually discontinued in successive days.
- If a permanent corticosteroid effect is required, oral therapy is recommended.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not handle by people hypersensitive to Dexamethasone.
- In case of contact with skin, eyes or mucous membranes, wash immediately with plenty of water.
- Do not handle by pregnant women.
- In case of accidental injection, go immediately to a medical center and show the product label.

ADVERSE EFFECTS

Adverse effects and adverse reactions:
May cause polyphagia, polyuria and polydipsia, delay wound healing, weaken resistance to infections or aggravate existing ones. In prolonged treatment it can cause hyperadrenocorticism iatrogenic (Cushing's syndrome), with loss of muscle mass, weakness, weight gain with fat redistribution, osteoporosis and ulcers.

GUARD PERIOD

Cattle, Sheep and Pigs:

- Meat: 5 days.
- Milk: 3 days.

Horses:

Do not administer to horses which meat is intended for human consumption.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Not Eliminate containers with product remains on the ground or water courses. For expired or not products used contact the manufacturer laboratory.

CONSERVATION

Store between 15 and 30 ° C, protected from light. Once the container is opened, use within 4 weeks. Discard the unused product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Ampule with 50 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

- Chile: Reg. SAG N °: 1366
- Uruguay: Reg. N° MGAP A-4495

HASYUN® ORAL TABLETS

COMPRIMIDO ORAL.

ANTIINFLAMATORIO Y ANTIALÉRGICO.

Technical Specification

SPECIES

Dogs and cats.

As a treatment in hypersensitive disorders such as state of shock; in immune disease treatments such as for Pemphigus; in serious cases of dermatitis and systemic lupus erythematosus; musculoskeletal disorders; allergies and hypoadrenocorticism.

DOSAGE FORM

Oral tablet .

THERAPEUTIC ACTION

Anti-inflammatory and anti-allergic.

COMPOSITION

Each tablet contains:
Dexamethasone.....025 mg
Excipients q.s.p.....1 tablet

INDICATIONS

- Do not use in pregnant females.
- Do not use in patients with gastric or duodenal ulcers.
- Do not use in animals with corneal ulcer.
- Do not use in patients with Diabetes Mellitus or Cushing's Syndrome.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route.

Product dose: Anti-inflammatory – Anti allergic:

- Dogs: 1 to 4 tablets (0.25 to 1 mg) daily, for 7 days
- Cats: ½ to 2 tablets (0.125 to 0.5 mg) daily, for 7 days.

DRUG INTERACTIONS

- Concomitant administration with Amphotericin B or caliuretic diuretics (Furosemide, thiazides) can cause hypokalemia.
- When both drugs are used together with digitalis glycosides, the possibility of toxicity can be increased if hypokalemia is generated.
- Dexamethasone can lower blood levels of salicylates.
- Insulin requirements may increase in patients receiving Dexamethasone.
- Phenytoin, Phenobarbital and Rifampicin can increase glucocorticoid metabolism.
- Concomitant administration with Cyclosporine can increase the blood levels of both drugs, with mutual inhibition of liver metabolism.
- Concomitant administration with ulcerogenic drugs (eg, non-steroidal anti-inflammatory drugs) may increase the risk of gastrointestinal ulceration.
- Patients treated with corticosteroids at immunosuppressive doses should not receive live attenuated virus vaccines, as they may enhance viral replication.

CONTRAINDICATIONS

- Do not use in pregnant females.
- Do not use in patients with gastric or duodenal ulcers.
- Do not use in animals with corneal ulcer.
- Do not use in patients with Diabetes Mellitus or Cushing's Syndrome.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Wash hands after administering the product.

WARNINGS

Advertencias y precauciones especiales de uso:

- Mantener fuera del alcance de los niños.
- En caso de terapia prolongada (más de 7 días) se debe suspender gradualmente en días sucesivos acorde a lo indicado por el Médico Veterinario tratante.

ADVERSE EFFECTS

Adverse effects and adverse reactions:
It can cause polyphagia, polyuria, and polydipsia, delay wound healing, weaken resistance to infections, or exacerbate existing infections. In prolonged treatments it can cause iatrogenic hyperadrenocorticism (Cushing's Syndrome), with loss of muscle mass, weakness, weight gain with fat redistribution, osteoporosis and ulcers.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
Do not dispose of containers with product residues on the ground or water courses. Empty containers can be discarded as household waste, without any special precautions. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Keep in a cool and dry place, at room temperature between 15 ° and 30° C.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Case with 10 tablets.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No.: 0527
- Uruguay: Reg. No. MGAP A-4494

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

HEMODRAG® 20 ML - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

HEMOSTÁTICO Y ANGIOPROTECTOR.

Technical Specification

SPECIES

Bovine, horses and pigs.

Hemodrag® is indicated for the prevention and treatment of bleeding in veterinary surgery, reproductive pathologies and obstetrics, accidental traumatology and localized bleeding in various organs. It has a hemostatic and angioprotective effect on skin processes such as petechiae, bruises and wounds.

Hemodrag® is effective for all bleeding processes whatever their etiology (with the exception of bleeding caused by coumarin agents), because it acts on primary haemostasis. It can be used in horse and bovine castration, tail docking and castration in pigs and in puerperal haemorrhage in cows and mares.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Hemostatic and Angioprotective.

COMPOSITION

Each ml contains:
Etamsilate.....125 mg
Excipients q.s.p.....1 mL

INDICATIONS

- Do not administer in cases of acute porphyriasis, since it has been seen to be porphyrogenic in animals.
- Do not use in animals with known hypersensitivity to the active substance.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration by slow intravenous or deep intramuscular route.

Dose of the active ingredient:
- Cattle, horses and pigs: 5 to 12.5 mg/Kg of weight, every six hours, for 1 to 3 days, as necessary.

Product dosage:
- Cattle, horses and pigs: 0.4 to 1 mL /10 kg of weight, every six hours, for 1 to 3 days, as necessary.

DRUG INTERACTIONS

Do not use concomitantly with other pharmacological products.

CONTRAINDICATIONS

- Do not administer in cases of acute porphyriasis, since it has been seen to be porphyrogenic in animals.
- Do not use in animals with known hypersensitivity to the active substance.

PRECAUTIONS

Special warnings and precautions for use

- Keep out of the reach of children and pets.
- This product is indicated for the treatment of animals individually. Do not treat the entire mass of animals.
- Animals must not be slaughtered for human consumption during and/or immediately after treatment.
- The product oxidizes in the presence of Oxygen, therefore the remaining volume after administration must be discarded.
- In the case of intramuscular administration, if the required dose is greater than 20 mL, it is suggested to divide it into two or more points of application.

SPECIAL PRECAUTIONS FOR USE

Special warnings and precautions for use

- Keep out of the reach of children and pets.
- This product is indicated for the treatment of animals individually. Do not treat the entire mass of animals.
- Animals must not be slaughtered for human consumption during and/or immediately after treatment.
- The product oxidizes in the presence of Oxygen, therefore the remaining volume after administration must be discarded.
- In the case of intramuscular administration, if the required dose is greater than 20 mL, it is suggested to divide it into two or more points of application.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Manage with caution. Wash hands after applying the product.
- People with known hypersensitivity to ethamsylate should avoid contact with the drug.
- Accidental self-injection may cause alteration of blood pressure. In case of occurrence, consult a doctor immediately and show the product label.

ADVERSE EFFECTS

They have not been described.

GUARD PERIOD

Meat: 0 days.

Milk: 0 days.

OBSERVATIONS

Overdose:
Studies carried out on the tolerance of ethamsylate indicate that it is well tolerated even in doses much higher than those recommended.

Special precautions for disposal of unused product or waste material:
Discard the remains of unused product in its original container. Do not throw the empty container or with remains of the product in rivers, lakes or streams of natural water. Dispose of this product carefully with your household waste.

CONSERVATION

Store at room temperature, between 15 and 30 °C, in a dry place protected from light.
Once the container is opened, use within 1 week. Discard unused product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Ampule with 20 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: SAG Reg. No. 1967
Peru: Reg. SENASA F.F3.01.I.0009
Imported and Distributed by Representaciones Durand SAC.
Av. Manuel Olguín N° 501 Office N° 604 Santiago de Surco Lima.
Costa Rica: Reg. No. MV-7150

COUNTRIES WHERE IT IS MARKETED

Imported and Distributed in Peru by:
Representations Durand SAC.
Av. Manuel Olguín N° 501 Office N° 604 Santiago de Surco Lima.

HEMODRAG® DOG 4 X 5 ML - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

HEMOSTÁTICO Y ANGIOPROTECTOR.

Technical Specification

SPECIES

Dogs.

Hemodrag® is indicated for the prevention and treatment of hemorrhages in veterinary surgery, reproductive and obstetric pathologies, accidental trauma and localized hemorrhages in various organs. It has a hemostatic and angioprotective effect on dermal processes such as petechiae, hematomas and wounds.

Hemodrag® is effective for all bleeding processes regardless of their etiology (with the exception of bleeding caused by coumarin agents), because it acts on primary hemostasis. In dogs it can be used for tail docking, castration and postpartum haemorrhages.

DOSAGE FORM

Injectable solution

THERAPEUTIC ACTION

Hemostatic and Angioprotective.

COMPOSITION

Each 1 mL contains:
Ethamsylate ..... 125 mg
Excipients q.s. ..... 1 mL

INDICATIONS

- Do not administer in cases of acute porphyriasis, since it has been seen to be porphyrogenic in animals.
- Do not use in animals with known hypersensitivity to the active principle.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration by slow intravenous or deep intramuscular route.

Dose of the active principle:
5 to 12.5 mg / Kg of weight, in a single dose.

Product dosage:
0.4 to 1 mL / 10 Kg of weight, in a single dose.

DRUG INTERACTIONS

Do not use concomitantly with other pharmacological products.

CONTRAINDICATIONS

- Do not administer in cases of acute porphyriasis, since it has been seen to be porphyrogenic in animals.
- Do not use in animals with known hypersensitivity to the active principle.

PRECAUTIONS

Special warnings and precautions for use

- Keep out of the reach of children.
- This product is indicated for the treatment of animals individually. Do not treat the entire mass of animals.
- Animals should not be slaughtered for human consumption during and / or immediately after treatment.
- The product oxidizes in the presence of oxygen, for which the remaining volume after administration must be discarded.
- In the case of intramuscular administration, if the required dose is greater than 20 mL, it is suggested to divide it into two or more application points.

SPECIAL PRECAUTIONS FOR USE

Special warnings and precautions for use

- Keep out of the reach of children.
- This product is indicated for the treatment of animals individually. Do not treat the entire mass of animals.
- Animals should not be slaughtered for human consumption during and / or immediately after treatment.
- The product oxidizes in the presence of oxygen, for which the remaining volume after administration must be discarded.
- In the case of intramuscular administration, if the required dose is greater than 20 mL, it is suggested to divide it into two or more application points.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Administer with caution. Wash your hands after applying the product.
- People with known hypersensitivity to ethamsylate should avoid contact with the drug.
- Accidental self-injection can cause alteration of blood pressure. In case of occurrence, immediately consult a doctor and show the product label.

OBSERVATIONS

Overdose:
Studies carried out on the tolerance of ethamsylate indicate that the drug is well tolerated even in doses much higher than those recommended.

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of natural water. Dispose of the waste of this product with care together with household waste.

CONSERVATION

Store at room temperature, between 15 ° C and 30 ° C, in a dry place protected from light. Once the container is opened, use within 1 week. Discard the unused product after that period of time.

PRESENTATION

4 vials of 5 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 1967
Perú: Reg. SENASA F.F3.01.I.0009

COUNTRIES WHERE IT IS MARKETED

Imported and Distributed in Peru by:
Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

HEPRO HORSE®

SUSPENSIÓN ORAL.

SUPLEMENTO NUTRICIONAL PARA CABALLOS.

Technical Specification

SPECIES

Horses.

DOSAGE FORM

Oral Supension

THERAPEUTIC ACTION

Nutritional Suplement for horses.

COMPOSITION

Each 100 mL of oral suspension contains:

Silymarin .....1.0 g

DL-Methionine.....1.6 g

Choline Chloride.....1.0 g

Vitamin B1 .....0.015 g

Vitamin B2 .....0,005 g

Vitamin B6 .....0.015 g

Vitamin B12 .....0.00028 g

Calcium Pantothenate .....0.01 g

Nicotinic Acid .....0.025 g

Excipients q.s.….....100 mL

PROPERTIES

HEPRO HORSE® is a nutritional supplement containing Silymarin, Methionine and B-Complex Vitamins, specially designed to nutritionally support and promote a healthy liver in horses.

Silymarin is a natural compound extracted from Milk Thistle (Silybum marianum), with powerful antioxidant and cytoprotective properties that help to stabilize and protect cell membranes. It also modulates liver metabolism, promoting its functioning and preventing toxic and drug aggression damages.

Due to its Methionine and B-Complex Vitamins, HEPRO HORSE® promotes adequate hepatic functions, stimulating energetic metabolism thus helping in horse health maintenance.

USE INSTRUCTIONS

Shake well before using.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer 100 mL per horse a day per oral route.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Keep in a cool and dry place protected from light, below 30 °C.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

1L o 3.8 L

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N°: RM 03-008N

NOTE

VETERINARY USE ONLY

EXCLUSIVE USE IN ANIMAL FEEDING

THIS PRODUCT IS NOT A COMPLETE FEED

HEPROTEC® EQUINE - SYRUP

JARABE.

SUPLEMENTO VITAMÍNICO. RECOMENDADO PARA MEJORAR EL RENDIMIENTO EN CABALLOS.

Technical Specification

SPECIES

Horses.

DOSAGE FORM

Syrup.

THERAPEUTIC ACTION

Vitamin supplement. Recommended to improve performance in horses.

COMPOSITION

Each 100 ml contains:

Methionine.....1 g

Vitamin B1 .....15 mg

Vitamin B2 .....5 mg

Vitamin B6 .....15 mg

Vitamin B12.....0.28 mg

Choline Chloride.....1 g

Calcium Pantothenate.....10 mg

Nicotinic acid.....25 mg

Excipients q.s.p.....100 mL

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route.

100 mL a day, during 5 days or according the professional guidance.

WARNINGS

Mantenga fuera del alcance de los niños.

OBSERVATIONS

This medicine does not contain liver extract.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° and 30°C.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Bottle with 500 ml and 3.8 liters.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 0941
- Dominican republic: Reg. No. 5968
- Panama: Reg. No. RF-3810-06

HEPROTEC® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

VITAMINAS, MINERALES, AMINOÁCIDOS.

Technical Specification

SPECIES

Bovine, horses , pigs, sheep, goats, dogs and cats.

Heprotec Injectable solution, is recommended as liver protector, hematopoietic. Especially recommended for pernicious and macrocytic anemia. Especially recommended for weak, convalescent or sick animals.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Vitamins, minerals, amino acids.

COMPOSITION

Each ampule contains:

Methionine..... 100 mg

Choline chloride..... 100 mg

Thiamine chloride..... 50 mg

Riboflavin 5.Sodium phosphate..... 7 mg

(Equivalent to 5.12 mg of Riboflavin base)

Nicotinamide..... 25 mg

Cyanocobalamin..... 100 µg

Excipients q.s.p..... 10 mL

ROUTE OD ADMINISTRATION AND DOSAGE

Administration route:

Intramuscular or intravenous route.

- Bovine, Horses: 2 ampules of 10 ml every 24 hours.
- Pigs, Sheep, Goats: 1 ampule of 10 ml every 24 hours.
- Dogs and Cats: 2 ml every 24 hours.

The duration of the treatment depends on the pathology and the general condition of the animal. It is recommended to prolong the treatment time when the animal is suffering from diarrhea or acute viral diseases.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

Fractional sale prohibited.

CONSERVATION

Store at room temperature between 15° and 30°C.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Box with 3 ampules of 10ml each.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 960
- Dominican Republic: Reg. No. 5605
- Panama: Reg. No. RF 3903-07

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

HEPROTEC® SYRUP

JARABE.

VITAMINAS DEL COMPLEJO B, MINERALES Y AMINOÁCIDO.

Technical Specification

SPECIES

Dogs.

Indicated as an appetite stimulant and adjuvant in the treatment of symptoms that present with deficiencies of some of these components, such as: liver deficiencies, toxicosis, antibiotic therapy, cirrhosis and jaundice.

DOSAGE FORM

Syrup.

THERAPEUTIC ACTION

Vitamin B complex, minerals and amino acids.

COMPOSITION

Each 100 ml contains:
Methionine.....1 g
Vitamin B1 .....15 mg
Vitamin B2 .....5 mg
Vitamin B6 .....15 mg
Vitamin B12.....0.28 mg
Choline chloride.....1 g
Calcium pantothenate .....10 mg
Nicotinic acid .....25 mg
Excipients q.s.p.....100 mL

INDICATIONS

Do not administer to animals with a history of hypersensitivity to any of the vitamins in the product.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route.

Dose:

- **Maintenance dose:** 1 tea spoon (5 mL) for each 10 Kg of weight, with the food, for 3 to 4 weeks.
- **As adjuvant dose:** 1 tea spoon (5 mL) for each 5Kg of weight, two or more times a day, for 3 to 4 weeks.

CONTRAINDICATIONS

Do not administer to animals with a history of hypersensitivity to any of the vitamins in the product.

SPECIAL PRECAUTIONS FOR THE OPERATOR

In case of skin contact, it is recommended to wash your hands with soap and plenty of water. If irritation develops and persists, see a doctor.
In the case of contact with the eyes, it is recommended to wash with plenty of water for 15 minutes. If irritation develops and persists, see a doctor.
In case of accidental ingestion, do not induce vomiting. Get medical help.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
Empty containers can be discarded as household waste, without any special precautions. Do not dispose of containers with product residues on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 ° and 30° C, protected from light. Once the container is opened, use within 5 weeks.
Discard the unused product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Bottle with 180 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- **Chile:** Reg. SAG. N° 941
- **Rep. Dominicana:** Reg. Nº 5968
- **Panamá:** Reg. N° RF-4129-18
- **El Salvador:** VE2013124887
- **Costa Rica:** Lic. DAA-MAG 579-025

COUNTRIES WHERE IT IS MARKETED

Distribution in El Salvador:
Rafael Alfredo Alfaro Castillo.
8a C. Pte. Pje. Moreno N ° 112, Col. Flor Blanca.
San Salvador, El Salvador.

Imported and distributed in Costa Rica by:
Proventas de Cartago S.R.L. 100 meters east Hogares Crea, San Blas. Carthage.
Tel: 2591 4624 Fax: 2591 5339

HERPLEX-L® ORAL SUSPENSION

SUSPENSIÓN ORAL.

SUPLEMENTO NUTRICIONAL.

Technical Specification

SPECIES

Cats.

Herplex-L® is a highly palatable, off-white to brown nutritional supplement specially formulated for cats.
It can be used in kittens from the first weeks of life.
L-Lysine is an essential amino acid in growing kittens, aiding in rapid weight gain. It favors the formation of collagen, fibrin and keratin, helping in the maintenance of the health of the skin and the connective tissue.

DOSAGE FORM

Oral suspension.

THERAPEUTIC ACTION

Nutritional supplement.

COMPOSITION

Each 1 mL of suspension contains:
L-lysine.....400 mg
Excipients q.s.p.....1 mL

INGREDIENTS

L-Lysine Hydrochloride, crab essence (flavoring) and authorized preservatives.

PROPERTIES

ROUTE OD ADMINISTRATION AND DOSAGE

Administer orally, 1.25 mL per cat, once daily for 25 to 30 days

WARNINGS

Almacenar fuera del alcance de los niños.

OBSERVATIONS

- It does not constitute complete food.
- Exclusive use in animal feed
- Includes dispenser.
- Flavored with crab meat.

CONSERVATION

Store between 15 ° and 30 ° C, protected from light. Do not refrigerate.

CONDITION OF SALE

Direct sales.

PRESENTATION

Bottle with 30 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. LENAA N°: RM 03-008N
Bolivia: Reg. SENASAG PUV-A nº 008927/19
Costa Rica: Lic. DAA-MAG 579-018

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

Imported and distributed in Costa Rica by:
Proventas de Cartago S.R.L.
100 meters east Hogares Crea, San Blas. Carthage.
Tel: 2591 4624 Fax: 2591 5339

INMUNOPET® ORAL SUSPENSION

SUSPENSIÓN ORAL.

ADITIVO A BASE DE BETAGLUCANOS.

Technical Specification

SPECIES

Dogs and cats.

Additive based on beta-glucans, specially formulated for dogs and cats.
Beta-glucans correspond to natural polysaccharides isolated from oyster mushroom *Pleurotus ostreatus*.

DOSAGE FORM

Oral suspension.

THERAPEUTIC ACTION

Betaglucan base additive

COMPOSITION

Each 100 mL of suspension contains:
Beta-(1,3/1,6)-D-glucans..... 1.0 g
Excipients q.s.p.....100 mL

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration. Betaglucans can be added to the food or administered directly to the oral cavity.

Recommended dose:
Dogs and cats:
Administer 2 mL for every 1 to 5 Kg of body weight, once a day for 2 to 3 months, or as needed.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

- Shake vigorously before use.
- It does not constitute a complete food.
- Exclusive use in animal feed

CONSERVATION

Store in a cool, dry place and away from light, at no more than 30ºC

CONDITION OF SALE

Over the counter.

PRESENTATION

Bottle with 60 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Reg. LENAA N°: RM 03-008N

INVEADE® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

VITAMINAS A, D, E.

Technical Specification

SPECIES

Bovine, sheep, goats and pigs.

- Promotes weight gain.
- Improves fertility.
- Improves the fur.
- Increases the defenses.
- Strengthening of the cow in the pre- and post-partum periods.
- Strengthening for Fall and Winter seasons.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

A,D, E Vitamins.

COMPOSITION

Each 1 mL contains:
Vitamin A.....500,000 U.I.
Vitamin D3.....75,000 U.I.
Vitamin E.....50 U.I.
Excipients q.s.p.....1 mL

ROUTE OD ADMINISTRATION AND DOSAGE

Bovine: intramuscular injection.

- **Calves:** 1 to1.5 mL/animal (Vitamin A 500,000-750,000 UI; Vitamin D3: 75,000-112.000 UI; Vitamin E: 50-75 UI).
- **Cows, heifers and steers:** 2-4 mL/animal (Vitamin A: 1,000,000-2,000,000 UI; Vitamin D3: 150,000-300,000 UI); Vitamin E: 100-200 UI).
- **Oxen and bulls:** 3.5-4 mL/animal (Vitamin A: 1,750,000-2,000,000 UI; Vitamin D3: 262,500-300,000 UI; Vitamin E: 175-200 UI).

Sheep and goats: Subcutaneous injection.

- **Lamb and ewe lambs:** 0.5-1 mL/animal (Vitamin A: 250,000-500,000 UI; Vitamin D3: 37,500-75,000 UI; Vitamin E: 25-50 UI).
- **Sheep and rams (jackets):** 0.5-1 mL/animal (Vitamin A: 250,000-500,000 UI; Vitamin D3: 37,500-75,000 UI; Vitamin E: 25-50 UI).
- **goatling:** 0.5-1 mL/animal (Vitamin A: 250,000-500,000 UI; Vitamin D3: 37,500-75,000 UI; Vitamin E: 25-50 UI).
- **Goats and snitches (jackets):** 0.5-1ml/animal (Vitamin A: 250,000-500,000 UI; Vitamin D3: 37,500-75,000 UI; Vitamin E: 25-50 UI).

Pigs: Intramuscular injection.

- **Piglets:** 0.5 mL (Vitamin A: 250,000 UI; Vitamin D3: 37,500 UI; Vitamin E: 25 UI).
- **Mother sows and boars:** 1-2 mL/animal (Vitamin A: 500,000-1,000,000 UI; Vitamin D3: 150,000-75,000 UI; Vitamin E: 50-100 UI).

The administration is in a single dose for all species.

SPECIAL PRECAUTIONS FOR THE OPERATOR

In case of contact with skin, eyes or mucous membranes, wash immediately with plenty of water.
If there is irritation, go immediately to a medical center and show the product label.
In case of accidental injection, go immediately to a medical center and show the product label.

WARNINGS

Almacenar fuera del alcance de los niños.

ADVERSE EFFECTS

Like all parenteral preparations, in exceptional cases it can produce an anaphylactic reaction.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 ° C, protected from light. Use immediately once opened and discard the excess product.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

10 mL, 50 mL and 100 mL vial

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No. 957

INVEADE® ORAL SOLUTION

SOLUCIÓN ORAL.

VITAMINAS A, D, E.

Technical Specification

SPECIES

Dogs.

- Growing puppies.
- Pregnant, lactating, senile animals.
- Hypovitaminosis A, D, E.
- Skin alteration associated to hypovitaminosis A and E.

DOSAGE FORM

Oral solution.

THERAPEUTIC ACTION

Vitamins A,D, E.

COMPOSITION

Each 1 mL contains:
Vitamin A Palmitate.....4,000 UI
Vitamin D2.....400 UI
Vitamin E Acetate.....22 UI
Excipients q.s.p.....1 mL

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route.

Product dose:

- Dogs, adults: 1 drop/2 Kg of weight / day during 1 month.
- Puppies and pregnant females: 1 drop/Kg of weight/day during 1 month.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- In case of contact with the skin, it is recommended to wash hands with soap and plenty of water. If irritation develops and persists, see doctor.
- In the case of contact with the eyes, it is recommended to wash with plenty of water for 15 minutes. If irritation develops and persists, see doctor.
- In case of accidental ingestion, do not induce vomiting. Get medical help.

WARNINGS

Almacenar fuera del alcance de los niños.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 ° C, protected from light.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Dropper bottle with 15 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG N° 950
Costa Rica: Lic. DAA-MAG 579-015

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Costa Rica by:
Proventas de Cartago S.R.L.
100 meters east Hogares Crea, San Blas. Carthage.
Tel: 2591 4624 Fax: 2591 5339

INVECLOR® - ANTISEPTIC SOAP

JABÓN LÍQUIDO. CLORHEXIDINA GLUCONATO 2%.

JABÓN ANTISÉPTICO DE PIEL Y MUCOSAS.

Technical Specification

SPECIES

Horses

INVECLOR®, Chlorhexidine Gluconate 2%, liquid soap, is indicated as an antiseptic soap for skin and mucous membranes.

DESCRIPTION

The INVECLOR® product combines the advantages of soapy cleaners with an effective, broad-spectrum and safe antiseptic such as Chlorhexidine, making it ideal for treating skin and / or mucous areas where it is required to reduce contamination quickly and persistent.

Chlorhexidine has a broad spectrum of antimicrobial activity. It is very effective against Gram positive bacteria, especially Staphylococcus aureus, other staphylococci, and various species of streptococci. It is less effective against Gram negatives such as Escherichia coli, Klebsiella, Salmonella, and Pseudomonas. It is capable of preventing the germination of bacterial spores. It is moderately active against *M. tuberculosis.* It is active against fungi such as *Candida albicans, Microsporum gypseum, M. canis and Trichophyton mentagrophytes* and yeasts. It has variable activity against viruses.

DOSAGE FORM

Liquid soap. Chlorhexidine Gluconate 2%.

THERAPEUTIC ACTION

Antiseptic soap for skin and mucous membranes.

COMPOSITION

Each 100 mL of soap contains:
Chlorhexidine Gluconate ..... 2 g
Excipients q.s.p ..... 100 mL

INDICATIONS

- Do not use in animals that are hypersensitive to chlorhexidine gluconate.
- Contraindicated in cases of animals with wounds, since its use delays the healing process.

MODE OF APPLICATION

Before applying this product, remove dirt and contamination from the area to be treated. Then, moisten the area and apply the product directly on the skin and / or mucous membranes in sufficient quantity so that the area is completely covered. Scrub (with a brush, brush, sponge or by hand) the area gently for 5 minutes.
Rinse with plenty of water to remove the entire product and dry with a towel, gauze or sterile compress.

ROUTE OD ADMINISTRATION AND DOSAGE

Topical use.

Place a large quantity of INVECLOR®, Chlorhexidine Gluconate 2%, liquid soap, on the skin and / or mucosa area to be treated so that it is completely covered by the product.

DRUG INTERACTIONS

Do not mix with other disinfectants and antiseptics.

CONTRAINDICATIONS

- Do not use in animals that are hypersensitive to chlorhexidine gluconate.
- Contraindicated in cases of animals with wounds, since its use delays the healing process.

PRECAUTIONS

- The applied product should be allowed to act for 5 minutes before rinsing.
- To improve antiseptic conditions, it is recommended to depilate the treated area before using this product.
- Do not use in the eyes, ears, or inside the animal's mouth; In case of contact wash with a lot of water.

SPECIAL PRECAUTIONS FOR USE

- The applied product should be allowed to act for 5 minutes before rinsing.
- To improve antiseptic conditions, it is recommended to depilate the treated area before using this product.
- Do not use in the eyes, ears, or inside the animal's mouth; In case of contact wash with a lot of water.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- It should not be handled by people who are hypersensitive to Chlorhexidine.
- Wash hands after use. The use of gloves is recommended.
- In case of contact with the skin or the eyes, it is recommended to wash with plenty of water. If irritation develops and persists, see doctor.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

Use during pregnancy and lactation:
Indicated for safe use in pregnant or lactating females.

CONSERVATION

Store in a dry place at temperatures between 2 and 30º C protected from light.

CONDITION OF SALE

Free sale.

VETERINARY USE.

EXTERNAL USE.

PRESENTATION

1 liter bottle

PREPARED BY

Laboratorio Drag Pharma.

RECORDS

Reg. SAG Nº 2242.

INVECLOR ® ANTISEPTIC SPRAY

SPRAY DE USO TÓPICO. CLORHEXIDINA GLUCONATO 0.5%.

SPRAY ANTISÉPTICO DE RÁPIDA ACCIÓN GERMICIDA CON EFECTO RESIDUAL

Technical Specification

SPECIES

WARNINGS

- Sólo para uso externo.
- Aplicar directamente sin diluir.
- Mantener fuera del alcance de los niños.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.
Lautaro N° 300 • Quilicura • Santiago • Chile
www.dragpharma.cl

RECORDS

Reg. SAG N° 1852

INVECTINA® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANTIPARASITARIO DE AMPLIO ESPECTRO

Technical Specification

SPECIES

Bovine, sheep, goat and pigs.

Endectocide indicated in the treatment of gastrointestinal parasites (*Ostertagia sp., Nematodirus sp., Oesophagostomum sp., Haemonchus sp., Cooperia sp., Strongylus sp., Trichuris sp., Ascaris sp. Trichostrongylus sp.*) And lung (*Dictyocaulus sp.*), including their migratory and inhibited larvae.
Effective in treating scabies mites (*Sarcoptes scabiei, Psoroptes sp.*), Lice (*Linognathus sp., Haematopinus sp.*) And *Oestrus ovis.*

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Endectocide Broad spectrum Antiparasitic

COMPOSITION

Each 1 mL of injectable solution contains:
Ivermectin.....10 mg
Excipients q.s.p.....1 mL

INDICATIONS

- Do not administer to animals younger than 4 weeks.
- Do not use in animals hypersensitive to Ivermectin

ACTION SPECTRUM

ROUTE OD ADMINISTRATION AND DOSAGE

Administration route:
Subcutaneous

Dosage and method of use:

- Cattle, sheep and goats: 1 mL / 50 Kg of weight (equivalent to 0.2 mg of Ivermectin / Kg of weight) in a single dose. Repeat the dose after 7 days in case of sheep scabies.
- Pigs: 3 mL / 100 Kg of weight (equivalent to 0.3 mg of Ivermectin / Kg of weight) in a single dose. Do not administer more than 10 mL per injection point.

DRUG INTERACTIONS

It is not recommended to use concomitantly with other pharmaceuticals.

CONTRAINDICATIONS

- Do not administer to animals younger than 4 weeks.
- Do not use in animals hypersensitive to Ivermectin

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not handle by people hypersensitive to Ivermectin
- In case of contact with skin, eyes or mucosa, wash immediately with plenty of water.
- Do not smoke, drink or eat while administering the product.
- In case of injection, immediately go to a medical center and show the product label.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

Inflammation may occur at the injection site that resolves without treatment.

GUARD PERIOD

Meat: 49 days.
Milk: Do not use milk from treated animals for human consumption.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
Discard the remains of unused product in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of water, since the product is extremely toxic to aquatic species. Do not throw empty containers or containers with the rest of the product together with household waste. Do not reuse the container.

CONSERVATION

Store between 15 and 30 ° C, protected from light. Once the container is opened, use within 12 weeks. Discard the unused product after that period of time.

CONDITION OF SALE

Sale under Veterinary Medical prescription.

PRESENTATION

Ampule bottle with 50 mL and 100 mL, bottle with 500 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG Nº 623-B
Bolivia: Reg. SENASAG PUV-F N° 007257/16
Perú: Registro SENASA F.09.01.I.0239

COUNTRIES WHERE IT IS MARKETED

Imported and Distributed in Bolivia by:
AGROGUARANI SRL
TEL: +(591)314-1401
Santa Cruz de la Sierra-Bolivia

Imported and Distributed in Peru by:
Representaciones Durand SAC.
Av. Manuel Olguín N° 501 Oficina N° 604 Santiago de Surco Lima.

INVECTINA® PLUS - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ENDECTOCIDA Y FASCIOLICIDA.

Technical Specification

SPECIES

Bovine, sheep and goats.

Indicated for the treatment and control of internal and external parasitism in cattle, sheep and goats.
Effective against mature and immature forms of gastrointestinal and lung nematodes, *Fasciola hepatica*, scabies, lice and *Oestrus ovis*.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Endectocide and fasciolicide.

COMPOSITION

Each 1 mL of injectable solution contains:
Ivermectin.....5 mg
Closantel.....125 mg
Excipients q.s.p.....1 mL

INDICATIONS

- Do not administer in animals less than 4 weeks of age.
- Do not administer to animals hypersensitive to avermectins, or to Closantel.

ACTION SPECTRUM

Bovine: *Ostertagia ostertagi, Dictyocaulus viviparus, Trichostrongylus axei, T. columbriformis, Nematodirus helvetianus, Haemonchus contortus, Trichuris spp. Oesophagostomum radiatum, Cooperia oncophora, C. punctata, Fasciola hepatica, Linognathus sp., Psoroptes ovis, Sarcoptes scabiei, Haematopinus sp.*

Sheep: *Ostertagia circumcincta, O. trifurcata, Trichostrongylus columbriformis, T. vitrinus, T. capricola, Nematodirus flicollis, N. spathiger, Haemonchus contortus, Cooperia curticei, Dictyocaulus sp., Fasciola hepatica, Linognathus sp., Oestrus ovis, Psoroptes sp., Sarcoptes scabiei.*

Goats: *Ostertagia sp., Trichostrongylus sp., Nematodirus sp., Haemonchus sp., Cooperia sp., Dictyocaulus sp., Fasciola hepatica, Linognathus sp., Oestrus ovis, Psoroptes sp., Sarcoptes scabiei.*

ROUTE OD ADMINISTRATION AND DOSAGE

Administration route: Subcutaneous, single dose.

Active ingredients dose:
0.2 mg/Kg Ivermectin and 5 mg/Kg de Closantel.

Product dose:
Bovine, sheep and goats: 1 ml/l25 kg of body weight, subcutaneous route, single dose.

CONTRAINDICATIONS

- Do not administer in animals less than 4 weeks of age.
- Do not administer to animals hypersensitive to avermectins, or to Closantel.

PRECAUTIONS

- In case of individual hypersensitivity, suspend treatment.
- Do not administer intramuscularly or intravenously.
- For heavier animals, the dose should be divided into different injection sites to avoid discomfort in the area.

SPECIAL PRECAUTIONS FOR USE

- In case of individual hypersensitivity, suspend treatment.
- Do not administer intramuscularly or intravenously.
- For heavier animals, the dose should be divided into different injection sites to avoid discomfort in the area.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not handle by people who are hypersensitive to Ivermectin or Closantel.
- Use of gloves when handling the product.
- In case of contact with skin, eyes or mucous membranes, wash immediately with plenty of water. If there is irritation, go immediately to a medical center and show the product label.
- In case of accidental injection, go immediately to a medical center and show the product label.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Meat: 49 days.

Milk: Do not use the milk of treated animals for human consumption.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 ° C, protected from light.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

100 mL vial and 500 mL vial

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 1375-B
- Dominican Republic: Reg. No. 5611

INVERMIC® CAT - ORAL SOLUTION

SOLUCIÓN ORAL.

ANTIPARASITARIO INTERNO.

Technical Specification

SPECIES

Cats.

- Anthelmintic against larval and adult forms of nematodes such as: *Toxascaris leonina, Toxocara cati, Ancylostoma spp. and Uncinaria stenocephala.*
- Immunostimulating action, which can be used in weakened or immunosuppressed animals.

DOSAGE FORM

Oral solution.

THERAPEUTIC ACTION

Internal antiparasitic.

COMPOSITION

Each 100 mL contains:
Levamisole Hydrochloride….....2.360 g
(Equivalent to 2.0 g of Levamisole)
Excipients q.s.p.....100 mL

INDICATIONS

- Do not administer in animals with hypersensitivity to the active principle.
- Do not administer to female cats in the last third of pregnancy or during lactation.
- Do not administer to cats with liver or kidney failure.
- Do not administer to cats under 4 weeks of age.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer directly into the oral cavity.

Dose:

- **Antiparasitic:** Administer 6 drops / kg of weight (6 mg / kg). Single dose.
- **Immunomodulator:** Administer 2 to 3 drops / Kg of weight (2 to 3 mg / Kg). Use 3 times a week.

The duration of treatment depends on the immune pathology to be treated.

CONTRAINDICATIONS

- Do not administer in animals with hypersensitivity to the active principle.
- Do not administer to female cats in the last third of pregnancy or during lactation.
- Do not administer to cats with liver or kidney failure.
- Do not administer to cats under 4 weeks of age.

WARNINGS

- Emplear con precaución al administrar en conjunto con otros compuestos nicotínicos o drogas inhibidoras de colinesterasa, pues pueden aumentar los efectos tóxicos del Levamisol.
- No utilizar en forma concomitante con Cloranfenicol
- Mantener fuera del alcance de los niños.

OBSERVATIONS

Specially flavored for the cat taste.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° and 30°C.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Dropper bottle with 10 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG No. 0368

INVERMIC® DOG - ORAL SOLUTION

SOLUCIÓN ORAL.

ANTIPARASITARIO INTERNO.

Technical Specification

SPECIES

Dogs.

- Anthelmintic against larval and adult forms of nematodes such as: *Toxascaris leonina, Toxocara canis, Ancylostoma caninum and Uncinaria stenocephala.*
- Immunostimulating action, which can be used in weakened or immunosuppressed animals.

DOSAGE FORM

Oral solution.

THERAPEUTIC ACTION

Internal Antiparasitic.

COMPOSITION

Each 100 mL contains.
Levamisole Hydrochloride.....2.360 g
(Equivalent to 2.0 g of Levamisole)
Excipients q.s.p.....100 mL

INDICATIONS

- Do not administer in animals with hypersensitivity to the active principle.
- Do not administer to female dogs in the last third of gestation or during the lactation period.
- Do not administer to dogs with liver or kidney failure.
- Do not administer to dogs under 2 weeks of age.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer directly into the oral cavity.

Dose:

- **Antiparasitic:** Administer 6 drops / kg of weight (6 mg / kg). Single dose.
- **Immunomodulator:** Administer 2 to 3 drops / Kg of weight (2 to 3 mg / Kg). Use 3 times a week.

The duration of treatment depends on the immune pathology to be treated.

CONTRAINDICATIONS

- Do not administer in animals with hypersensitivity to the active principle.
- Do not administer to female dogs in the last third of gestation or during the lactation period.
- Do not administer to dogs with liver or kidney failure.
- Do not administer to dogs under 2 weeks of age.

WARNINGS

- Emplear con precaución al administrar en conjunto con otros compuestos nicotínicos o drogas inhibidoras de colinesterasa, pues pueden aumentar los efectos tóxicos del Levamisol.
- No utilizar en forma concomitante con Cloranfenicol.
- Mantener fuera del alcance de los niños.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° and 30°C.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Dropper bottle with 10 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile:Reg. SAG No. 0368

INVERMIC® PIG - ORAL POWDER

POLVO ORAL.

ANTIPARASITARIO INTERNO.

Technical Specification

SPECIES

Pigs.

Internal antiparasitic effective against the adult and larval stages of gastrointestinal nematodes (*Ascaris suum, Oesophagostomum spp. And Strongyloides ransomi*) and lungs (*Stephanurus dentatus* (present on Easter Island) and *Metastrongylus spp*.) In pigs.

DOSAGE FORM

Oral powder.

THERAPEUTIC ACTION

Internal antiparasitic.

COMPOSITION

Each 100 g contains:
Levamisole Hydrochloride.....5.5 g
(Equivalent to 4.6 g of Levamisole as free base)
Excipients q.s.p.....100 g

INDICATIONS

- Do not administer to animals hypersensitive to Levamisole.
- Do not administer in the last third of pregnancy, lactation or in reproductive animals.

USE INSTRUCTIONS

Recommendations for incorporating Invermic® 5.5% into drinking water:

1. Add the amount of Invermic® 5.5% according to the number of animals to be treated and dissolve in a fraction of water to be administered, then complete the total of the indicated volume, which must correspond to one third (1/3) of the total estimated daily consumption according to age, weight and weather conditions.

2. Suspend the administration of water through automatic drinkers, while the administration of Invermic® 5.5% lasts.

3. The preparation and administration of the medicine must be carried out in troughs intended only for the treatment of sick animals.

4. It is recommended to suspend the consumption of water a couple of hours before the administration of the product dissolved in the water.

5. The medicine should be administered in a single daily dose. Do not reuse remnants of prepared solution.

6. Prepare solutions daily. The solution must be prepared under the supervision of a Veterinarian.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral route, in drinking water.

Dose of the active principle:
8 mg / Kg of live weight.

Product dosage:
Dissolve 20 g of product with water, for 115 kg of live weight.

DRUG INTERACTIONS

Do not use concomitantly with other Nicotinic compounds (eg Pyrantel, Morantel, Diethylcarbamazine) or cholinesterase inhibitors (eg Organophosphates, Neostigmine).

CONTRAINDICATIONS

- Do not administer to animals hypersensitive to Levamisole.
- Do not administer in the last third of pregnancy, lactation or in reproductive animals.

PRECAUTIONS

- Use with caution in animals that are very weakened or show significant kidney or liver damage.
- Delay use in animals stressed by vaccination or castration.

SPECIAL PRECAUTIONS FOR USE

- Use with caution in animals that are very weakened or show significant kidney or liver damage.
- Delay use in animals stressed by vaccination or castration.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not handle by people hypersensitive to Levamisole.
- Do not handle by pregnant women.
- In case of contact with skin, eyes or mucous membranes, wash immediately with plenty of water.
- In case of accidental ingestion, go immediately to a medical center and show the product label.

WARNINGS

Almacenar fuera del alcance de los niños.

ADVERSE EFFECTS

Coughing, salivation, foaming at the mouth, mild vomiting, abdominal pain, and diarrhea may be observed transiently.

GUARD PERIOD

Meat: 28 days.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 ° C, protected from light.
Once opened or prepared, use immediately and discard the excess product.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Case with 25 sachets of 20 g each.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG N° 573
- Panamá: Reg. N° RF-4135-18

INVESPAS® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANTIDIARREICO PARA BOVINOS, OVINOS Y CERDOS

Technical Specification

SPECIES

Bovine, sheep and pigs.

Supportive therapy in the treatment of diarrhea in cattle, sheep and pigs.

Nutritional diarrhea: The best results are achieved when the conditions allow to combine Invespas® treatment, with the necessary dietary changes.

Infectious diarrhea: Invespas® is an antidiarrheal and antispasmodic. When the infectious therapy is combined with Invespas® used as symptomatic treatment, the diarrhea stop faster and as a result the dehydration is limited.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Antidiarrheal for cattle, sheep and pigs

COMPOSITION

Each mL contains:
Benzetimide Hydrochloride.....0.165 mg
(Equivalent to 0.15 mg de Benzetimide)
Excipients q.s.p.....1 mL

INDICATIONS

Do not administer Invespas® to pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Intramuscular administration.

Bovine and sheep:
15 µg de Benzetimide for of weight or 1 mL each 10 Kg of weight, with a maximum of 20 mL per animal.

Pigs:
60 µg of Benzetimide for Kg of weight or 1 mL every 2.5 Kg of weight.

If is it needed, the Invespas® treatment can be repeated after 24 hours.

CONTRAINDICATIONS

Do not administer Invespas® to pregnant or lactating females.

WARNINGS

Mantener fuera del alcance de los niños.

SIDE EFFECTS

At therapy dose, occasionally can be observed anticholinergic effects such as tachycardia, mydriasis, tympanites (bloat) and bronchodilation.

GUARD PERIOD

Meat: 0 days.
Milk: 0 days.

CONSERVATION

- Store in a cool and dry place, at room temperature between 15º and 30º C.
- Check the expiration date on the bottle.

CONDITION OF SALE

To be supply only with veterinary prescription.

PRESENTATION

Ampule bottle with 50 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No. 949

INVETROID® EMULSIFIABLE SOLUTION

SOLUCIÓN EMULSIFICABLE.

ANTIPARASITARIO EXTERNO DE EFECTO RESIDUAL.

Technical Specification

SPECIES

Bovine, sheep and pigs.

Invetroid® is an effective antiparasitic in the control of lice (Bovicola ovis, Bovicola bovis, Linognathus pedalis, Linognathus ovillus, Linognathus vituli, Haematopinus eurysternus, Haematopinus suis), scabies mites (Psoroptes bovis, Psoroptes bobiei, Psoroptes bobiei var. , Sarcoptes scabiei var. Suis) and melophagi (Melophagus ovinus) in cattle, sheep and pigs.

DOSAGE FORM

Emulsifiable solution.

THERAPEUTIC ACTION

External antiparasitic with residual effect.

COMPOSITION

Each 100 mL of concentrated solution contains:
Cypermethrin..... 20 g
Excipients q.s.p.....100 mL

INDICATIONS

- Do not bathe in hours of intense heat, or when the animals are very tired and thirsty.
- Do not use during advanced stages of pregnancy or during lactation.
- Do not administer to animals with hypersensitivity to cypermethrin.

ROUTE OD ADMINISTRATION AND DOSAGE

Cattle and Sheep:
- Baths for scabies and lice: Dilute 1 liter of Invetroid® in 1,000 liters of water (equivalent to a final bath concentration of 0.02%). For replacement and reinforcement, use a dilution of 1.5 liters of the product in 1,000 liters of water (equivalent to a concentration of 0.03%); This is when the bathtub level has dropped by around 10%.

- For dry replacement and reinforcement: Dilute 500 mL of Invetroid® in 1,000 liters of water (equivalent to a final concentration of 0.01%). This dilution can be used as a spray, applying at a rate of 3-4 liters per animal.

Sheep Melophagus (Melophagus ovinus: "false tick"):

- Immersion bath: Loading: 1 liter of Invetroid® in 10,000 liters of water (equivalent to a final bath concentration of 0.002%). Replacement: 1 liter in 8,000 liters of water (equivalent to a concentration of 0.025%), when the level has dropped by 20%. It must be ensured that the animals are well wet, totally submerging them at least twice. Every 10,000 animals the bathroom must be renewed. Once the job is finished, all the remaining liquid must be removed.

- Aspersion: 1 liter of Invetroid® in 5,000 liters of water (equivalent to a final concentration of 0.004%). This system is recommended as a way to lower the melophage load in freshly shorn animals.

Pigs:
- Scabies and lice: Aspersion: Dilute 1 mL of Invetroid® in 5 liters of water (equivalent to a final concentration of 0.004%). Apply at a rate of 3-4 liters per animal. Apply using a spray pump, distributing the product evenly over the animal.

Frequency and duration of treatment: Single dose.

DRUG INTERACTIONS

Do not use concomitantly with other pharmaceutical products.

CONTRAINDICATIONS

- Do not bathe in hours of intense heat, or when the animals are very tired and thirsty.
- Do not use during advanced stages of pregnancy or during lactation.
- Do not administer to animals with hypersensitivity to cypermethrin.

PRECAUTIONS

- It must be ensured that both the immersion baths and the spray pumps are completely free of residues of other products that may have been applied previously.
- Use the mixture in a well-ventilated environment, avoiding contact with the animal's eyes.
- In case of rain threat, suspend the bath. In case of rain within 24 hours after bathing, repeat the treatment.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR USE

- It must be ensured that both the immersion baths and the spray pumps are completely free of residues of other products that may have been applied previously.
- Use the mixture in a well-ventilated environment, avoiding contact with the animal's eyes.
- In case of rain threat, suspend the bath. In case of rain within 24 hours after bathing, repeat the treatment.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- The applicator must wear a mask and gloves as a protection element for handling the product.
- Avoid contact with the eyes, skin and mucous membranes, Cypermethrin is an irritant and can cause skin sensitization.
- Contact with treated animals should be avoided until they are completely dry.
- Wash contaminated clothing before reuse.
- In case of contact with the skin, wash with plenty of water. In case of eye contact, wash immediately with plenty of water for at least 15 minutes.

ADVERSE EFFECTS

In animals with hypersensitivity to cypermethrin, dermal irritation may be observed.

GUARD PERIOD

Meat:

- Bovine: 14 days
- Sheep and pigs: 7 days.

Milk: 7 days.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Do not dispose of the empty container and / or product remains on the ground, rivers, lakes or streams of water. Any unused medicine or waste derived from it must be disposed of in an environmentally safe way.

CONSERVATION

Store below 30°C. Do not refrigerate or freeze.
Use immediately after opening and discard the excess product.
Use immediately once the solution is prepared and discard the excess product.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Bottle with 100 ml and 1 L.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG N° 955
Rep. Dominicana: Nº 9685

ITRASKIN® ORAL SOLUTION

SUSPENSIÓN ORAL.

ANTIFÚNGICO.

Technical Specification

SPECIES

Dogs and cats.

Itraskin®, oral suspension, is recommended for the treatment of fungal infections caused by microorganisms susceptible to Itraconazole, such as skin dermatophytosis caused by *Microsporum canis* in cats, dermatitis caused by *Malassezia pachydermatis* in dogs and systemic infections of fungal origin, such as, histoplasmosis, cryptococcosis, blastomycosis and sporotrichosis.

DOSAGE FORM

Oral suspension.

THERAPEUTIC ACTION

Antifungal.

COMPOSITION

Each mL of suspension contains:
Itraconazole.....20 mg
Excipients q.s.p.....1 mL

PROPERTIES

Itraconazole is a synthetic antifungal agent, belonging to the family of azole derivatives. It was introduced in the 90s in order to broaden the antifungal spectrum, increase its potency compared to other azoles and reduce adverse effects at the digestive level, especially in prolonged therapies.
The antifungal action of Itraconazole is generated through a multiple mechanism initiated by the inhibition of two cytochromes P450 involved in the biosynthesis of Ergosterol: CYP51 (sterol-14 alpha demethylase) and CYP61 (Delta 22-denaturase). Ergosterol is an essential component of the plasma membrane of fungi.

INDICATIONS

- Do not use in patients with hypersensitivity to itraconazole or other azole agents.
- Do not use in patients with hepatic impairment or hypochlorhydria.
- Do not use in females during the gestation and lactation period.
- Do not use in dogs and cats under 2 months of age.

ROUTE OD ADMINISTRATION AND DOSAGE

Orally.

Recommended dose:

Cats:

Pathology	Dose
Dermatophytosis by Microsporum canis:	1 mL every 2 Kg of weight (equivalent to 10 mg/kg) every 24 hours, for 4 weeks. It is recommended to continue the therapy until mycological cure or lack of dermatophyte isolation in three consecutive weekly cultures.
Systemic fungal infections (histoplasmosis, cryptococcosis, blastomycosis, etc.):	0.5 mL every 2 of weight (equivalent to 5 mg/kg) every 12 hours, or 1 ml every 2 Kg of weight (equivalent to 10 mg/kg) every 24 hours. The therapy should be continued for 2 to 3 months or until 1 month after the resolution of the clinical signs of the animal.

Dogs:

Patología	Dosis
Dermatitis por *Malassezia pachydermatis*:	2,5 mL cada 10 kg de peso (equivalentes a 5 mg/kg) cada 24 horas, por 3 semanas.
Enfermedades fúngicas sistémicas (blastomicosis, histoplasmosis, esporotricosis, etc.):	2,5 mL cada 10 kg de peso (equivalentes a 5 mg/kg) cada 24 horas, por 2 a 3 meses, o hasta después de 1 mes de la resolución de los signos clínicos del animal.

DRUG INTERACTIONS

- Itraconazole absorption is reduced by administration in conjunction with proton pump inhibitor antacids (Omeprazole) and H2 blockers (Cimetidine, Ranitidine, etc.) or Didanosine.
- Itraconazole can increase prothrombin times in patients receiving Warfarin or other coumarin anticoagulants. Rifampin can increase the metabolism of Itraconazole.
- Itraconazole can increase the serum levels of oral antidiabetic agents (eg Chlorpropamide, Glipizide etc.), which can lead to hypoglycemia.
- Do not use in conjunction with Cisapride, as it can cause ventricular arrhythmias.
- Itraconazole interacts with certain antihistamines (Ternadine, Astemizole), Benzodiazepines, calcium channel blockers, anticonvulsants, some antimicrobials and Cyclosporine.
- It is not recommended to use concomitantly with anticholinergics.

CONTRAINDICATIONS

- Do not use in patients with hypersensitivity to itraconazole or other azole agents.
- Do not use in patients with hepatic impairment or hypochlorhydria.
- Do not use in females during the gestation and lactation period.
- Do not use in dogs and cats under 2 months of age.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Wash your hands after administering the product.
- In the case of contact with the eyes, it is recommended to wash with plenty of water for 15 minutes.
- In case of accidental ingestion do not induce vomiting. Obtain medical attention if necessary.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

In dogs, the most common adverse effect is anorexia, especially with higher doses.Some cases may show signs of hepatotoxicity, in which case treatment should be discontinued at least temporarily. Liver damage is determined by increased ALT activity. Anorexia is often a symptomatic marker of toxicity and usually occurs in the second month of treatment. Some dogs treated with doses of 10 mg / kg may develop vasculitis or ulcerative skin lesions and edema of the limbs that may require a reduction of the dose to 5 mg / kg. These problems usually resolve after stopping the medication. In cats, adverse effects are dose related. Gastrointestinal effects (anorexia, weight loss, vomiting), hepatotoxicity (increased ALT, jaundice), and depression have been reported. If adverse effects occur and ALT is elevated, the drug should be discontinued. Enzyme hyperactivity in the absence of other signs does not necessarily require reducing the dose or stopping the medication. Once ALT levels normalize and other adverse effects decrease, and if necessary, the drug can be restarted at half the dose initially used with closer clinical supervision.

CONSERVATION

- Store at room temperature between 15 ° and 30 ° C.
- Once opened, use the product within 3 months. Discard the unused product after that period of time.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

Battle with 120 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 2053
Costa Rica: Reg. N° MAG CL4-15-13-5639
Rep. Dominicana: Reg. Nº 9305
Bolivia: Reg. SENASAG PUV-F N°007252/16

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

Imported and Distributed in Peru by:

Representaciones Durand SAC.
Av. Manuel Olguín N ° 501 Office N ° 604
Santiago de Surco Lima.

KAUPOL® BOVINE - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

VITAMINAS, GLUCOSA Y MINERALES.

Technical Specification

SPECIES

Bovine, pigs, sheep and goats.

Indicated in dehydration and in glucose and mineral deficiencies, which can be caused by: bleeding, diarrhea, burns, postpartum deficiencies and / or poisoning.

In production animals Kaupol®, administered concomitantly with vitamins B_1, B_6 and B_{12}, is indicated as a hepatic, lipotropic and hematopoietic protector.

Especially recommended for animals that are weakened, convalescing, sick or subjected to long demands.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Vitamins, Glucose and Minerals.

COMPOSITION

100 mL of injectable solution contains:
Riboflavin 5 Sodium phosphate.....10 mg
Nicotinamide.....200 mg
Anhydrous D-glucose (Dextrose).....10.0 g
Sodium.....11.96 mEq
Potassium.....0.54 mEq
Magnesium.....0.049 mEq
Calcium.....0.163 mEq
Chloride.....12.92 mEq
Excipients q.s.p.....100 mL

INDICATIONS

Do not administer in pathological states that require low sodium intake.

ROUTE OD ADMINISTRATION AND DOSAGE

Slow and subcutaneous intravenous administration.

Product dosage:

- Cattle and horses (greater than or equal to 400 kilos): 1000 - 2000 mL via E.V.
- Cattle (greater than or equal to 400 kilos): 200 - 300 mL via S.C.
- Cattle and Horses (200 - 300 kilos): 500 mL via E.V.
- Cattle (200 - 300 kilos): 50 to 80 mL via S.C.
- Sheep or Goats (25 - 60 kilos): 100 - 250 mL via E.V.
- Sheep or Goats (25 - 60 kilos): 40 mL via S.C. Pigs (greater than or equal to 120 kilos): 300 - 500 mL via E.V.
- Pigs (greater than or equal to 120 kilos): 300 - 500 mL via E.V. (Indicated in dehydration and bleeding).

CONTRAINDICATIONS

Do not administer in pathological states that require low sodium intake.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Meat: 0 days.

Milk: 0 dads.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° and 30°C., protected from sunlight.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

500 mL container

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 1351
- Costa Rica: Reg. No. MAG CL4-19-01-4140
- Dominican republic: Reg. No. 5914
- Panamá: Reg. N° RF-4133-18

KAUPOL® EQUINO - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

VITAMINAS, GLUCOSA Y MINERALES.

Technical Specification

SPECIES

Horses.

Indicated in dehydration and in glucose and mineral deficiencies, which can be caused by: bleeding, diarrhea, burns, postpartum deficiencies and / or poisoning.

In production animals Kaupol®, administered concomitantly with vitamins B_1, B_6 and B_{12}, is indicated as a hepatic, lipotropic and hematopoietic protector.

Especially recommended for animals that are weakened, convalescing, sick or subjected to long demands.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Vitamin, Glucose and Minerals.

COMPOSITION

100 mL of injectable solution contains:
Riboflavin 5 Sodium phosphate.....10 mg
Nicotinamide.....200 mg
Anhydrous D-glucose (Dextrose).....10.0 g
Sodium.....11.96 mEq
Potassium.....0.54 mEq
Magnesium.....0.049 mEq
Calcium.....0.163 mEq
Chloride.....12.92 mEq
Excipients q.s.p.....100 mL

INDICATIONS

Do not administer tin pathological states that require low sodium intake.

ROUTE OD ADMINISTRATION AND DOSAGE

Slow intravenously administration.

Product dose:

- Horses (weighing more than or equal to 400 Kg): 1000 - 2000 mL I.V. route.
- Horses (200 - 300 Kg): 500 mL I.V. route.

CONTRAINDICATIONS

Do not administer tin pathological states that require low sodium intake.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Meat: 0 day.

Milk: 0 day.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° and 30°C, protected from sunlight.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

500 mL container

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 1351
- Costa Rica: Reg. No. MAG CL4-19-01-4140
- Dominican Republic: Reg. No. 5914

KET-10® BOVINE - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANALGÉSICO, ANTIPIRÉTICO Y ANTIINFLAMATORIO NO ESTEROIDAL

Technical Specification

SPECIES

Bovine and pigs.

In cattle, indicated as an anti-inflammatory, analgesic and antipyretic treatment, particularly in:
respiratory conditions, acute bacterial mastitis, calf hyperthermia and in musculoskeletal conditions such as lameness, arthritis, trauma.
In pigs, indicated in the relief of pain, fever and inflammation in respiratory processes and in metritis syndrome, mastitis, agalactia, in conjunction with antibiotics.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

ANALGESIC, ANTIPYRETIC AND NON-STEROIDAL ANTI-INFLAMMATORY

COMPOSITION

Each mL of injectable solution contains:
Ketoprofen..... 100 mg
Excipients q.s.p.....1 mL

INDICATIONS

- Do not administer in pregnant females.
- Do not use in cases of severe kidney failure.
- Do not use simultaneously with diuretics or anticoagulants.
- Do not use in animals with heart or liver diseases, gastrointestinal ulcers or blood dyscrasias.
- Do not use in animals with known hypersensitivity to the active substance.

ROUTE OD ADMINISTRATION AND DOSAGE

Bovine

Administration route: Intramuscular and intravenous

Dose: 3 mg of Ketoprofen / Kg of live weight, equivalent to 3 mL of product / 100 Kg of weight.

Pigs
Administration route: Intramuscular.

Dose: 3 mg of Ketoprofen / Kg of live weight, equivalent to 3 mL of product / 100 Kg of weight.

DRUG INTERACTIONS

Simultaneous use with other non-steroidal anti-inflammatory drugs can increase side effects.

CONTRAINDICATIONS

- Do not administer in pregnant females.
- Do not use in cases of severe kidney failure.
- Do not use simultaneously with diuretics or anticoagulants.
- Do not use in animals with heart or liver diseases, gastrointestinal ulcers or blood dyscrasias.
- Do not use in animals with known hypersensitivity to the active substance.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Meat: 4 days.

Milk: 0 days

CONSERVATION

Store at a temperature between 2 ° and 30 ° C, protected from light. Once opened, use the product within 3 months. Discard the excess product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

50 mL vial

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG N° 1472
Bolivia: Reg. SENASAG PUV-F N° 007258/16
Perú: Registro SENASA F.99.01.I.0069

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:
AGROGUARANI SRL
TEL: + (591) 314-1401
Santa Cruz de la Sierra, Bolivia

Imported and Distributed in Peru by Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

KET-10® EQUINE - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANTIINFLAMATORIO NO ESTEROIDAL

Technical Specification

SPECIES

Horses.

In horses, indicated for the relief of inflammation and pain associated with musculoskeletal osteoarticular alterations, such as lameness of traumatic origin, arthritis, osteoarthritis, tendonitis, bursitis, post-surgical inflammations, among others. It is also indicated in the symptomatic treatment of equine colic and in diseases of the respiratory system that involve inflammatory and painful conditions.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Non-steroidal anti-inflammatory

COMPOSITION

Each mL of injectable solution contains:
Ketoprofen.....100 mg
Excipients q.s.p.....1 mL

INDICATIONS

- Do not administer in pregnant females.
- Do not use in cases of severe kidney failure.
- Do not use simultaneously with diuretics or anticoagulants.
- Do not administer to foals under 2 weeks of age.
- Do not use in animals with heart or liver diseases, gastrointestinal ulcers or blood dyscrasias.
- Do not use in animals with known hypersensitivity to the active substance.

ROUTE OD ADMINISTRATION AND DOSAGE

Intravenous administration.

Dose of the active principle: 2 - 3 mg / Kg of weight per day.

Product dose: 1 mL for every 45 kilos of weight per day.

DRUG INTERACTIONS

Simultaneous use with other non-steroidal anti-inflammatory drugs can increase side effects.

CONTRAINDICATIONS

- Do not administer in pregnant females.
- Do not use in cases of severe kidney failure.
- Do not use simultaneously with diuretics or anticoagulants.
- Do not administer to foals under 2 weeks of age.
- Do not use in animals with heart or liver diseases, gastrointestinal ulcers or blood dyscrasias.
- Do not use in animals with known hypersensitivity to the active substance.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Do not use treated horse meat for human consumption.

CONSERVATION

Store at a temperature between 2 ° and 30 ° C, protected from light. Once opened, use the product within 3 months. Discard the excess product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

50 mL vial

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG Nº 1472
- Bolivia: Reg. SENASAG PUV-F N° 007258/16
- Perú: Reg. SENASA F.99.01.I.0069

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:
AGROGUARANI SRL
TEL: + (591) 314-1401
Santa Cruz de la Sierra, Bolivia

Imported and Distributed in Peru by Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

KETOSTOP® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANESTÉSICO.

Technical Specification

SPECIES

Dogs, cats and horses.

Ketostop® is a short-acting, fast induction, dissociative anesthetic. Recommended for emergency surgeries, short-term interventions such as caesarean sections, dystocic deliveries and surgeries that do not require relaxation of the skeletal muscles, causing immediate superficial analgesia and anesthesia without bulbar depression.

Ketostop® is characterized by producing a dissociative state, where the patient is observed outside his environment, determining during induction an apparently superficial but insensitive anesthesia to surgical stimulation and later amnesia, inducing a cataleptoid state. Upon awakening there may be hallucinations depending on external excitations.

DOSAGE FORM

Injectable solution

THERAPEUTIC ACTION

Anesthetic.

COMPOSITION

Each mL of the solution contains:
Ketamine Hydrochloride .....115.36 mg
(Equivalent to 100 mg of Ketamine base)
Excipients q.s.p.....1 mL

PROPERTIES

INDICATIONS

- Only anesthesia in horses, donkeys and dogs.
- Not recommended as a single agent in cesarean section.
- Not recommended as the sole agent of anesthesia for abdominal surgery or orthopedics.
- Renal and/or hepatic damage.
- Do not administer in pregnant or lactating females.
- High blood pressure, hypertension and severe intracranial hypertension.
- Severe cardiac decompensation.
- Animals under organophosphates treatment.

ROUTE OD ADMINISTRATION AND DOSAGE

For horses, dogs and cats, the following doses are recommended:

Active ingredient dose:
11 - 20 mg/Kg IM route
2 - 8 mg/Kg IV. route

Product dose:
0.11 – 0.2 ml/Kg IM route.
0.02 – 0.08 ml/Kg IV route.

DRUG INTERACTIONS

- Narcotics, barbiturates, or Diazepam can prolong recovery time after Ketamine anesthesia.
- When used with Halothane, recovery from Ketamine can be prolonged and its cardio-stimulating effects can be inhibited. Close supervision is recommended when using Ketamine with Halothane.
- Chloramphenicol (parenteral) can prolong the anesthetic actions of Ketamine.
- Neuromuscular blockers (for example Succinylcholine or Tubocurarine) can increase or prolong respiratory depression.

CONTRAINDICATIONS

- Only anesthesia in horses, donkeys and dogs.
- Not recommended as a single agent in cesarean section.
- Not recommended as the sole agent of anesthesia for abdominal surgery or orthopedics.
- Renal and/or hepatic damage.
- Do not administer in pregnant or lactating females.
- High blood pressure, hypertension and severe intracranial hypertension.
- Severe cardiac decompensation.
- Animals under organophosphates treatment.

PRECAUTIONS

- The highest doses correspond to the smaller species.
- The dosage calculated from the normal dose may vary depending on the patient, the intervention, the premedication and its effect.
- In dogs and horses, premedication is highly recommended.
- Cats' eyes are kept open after receiving Ketamine and should be protected with an ophthalmic lubricant to avoid excessive desiccation of the cornea.
- To minimize the incidence of reactions, it is recommended to reduce exposure to handling or loud noises during the recovery period.

SPECIAL PRECAUTIONS FOR USE

- The highest doses correspond to the smaller species.
- The dosage calculated from the normal dose may vary depending on the patient, the intervention, the premedication and its effect.
- In dogs and horses, premedication is highly recommended.
- Cats' eyes are kept open after receiving Ketamine and should be protected with an ophthalmic lubricant to avoid excessive desiccation of the cornea.
- To minimize the incidence of reactions, it is recommended to reduce exposure to handling or loud noises during the recovery period.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Wash hands after administering the product.

WARNINGS

Mantener fuera del alcance de los niños.

SIDE EFFECTS

ADVERSE EFFECTS

- There may be transient pain at the injection site when administered intramuscularly.
- After the administration of Ketamine, in most of the treated animals, the eyes remain open and must be protected with ophthalmic lubricants.
- Ketamine administration increases cardiac volume, heart rate and blood pressure in treated animals.
- With the higher recommended doses, there may be respiratory depression, vomiting, vocalizations, erratic and prolonged recovery, dyspnea, spastic movements, muscle tremors, hypertonicity, tonic / clonic seizures, opisthotonos and cardiac arrest. Muscle disorders and seizures can be controlled by administering Accepromazine or short-acting barbiturates.
- When used as a sole anesthetic, Ketamine can induce seizures in dogs and up to 20% of cats. The incidence of seizures is reduced when combined with Diazepam or Midazolam.
- Ketamine use can cause hypersalivation. Atropine premedication can reduce this effect.

GUARD PERIOD

Do not use treated horse meat for human consumption.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Dispose of this product waste carefully with household waste. Contact the manufacturing company or companies specialized in waste disposal, to receive recommendations on the disposal of expired or unused products.

CONSERVATION

Store in a cool place, protected from light, at room temperature between 15 and 25° C.

CONDITION OF SALE

Sale held under veterinary prescription controlled balance.

PRESENTATION

50 mL vial

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No. 0558

LAGRIPET® EXTERNAL SOLUTION

SOLUCIÓN EXTERNA.

LIMPIADOR DE MANCHAS CAUSADAS POR EL LAGRIMEO.

Technical Specification

SPECIES

Dogs and cats.

Lagripet® is a product especially formulated to clean and eliminate yellow or brown-yellowish colored stains, produced in the coat under the eyes due to tears. Its frequent use prevents the coat discoloration caused by tears.

DOSAGE FORM

External solution.

THERAPEUTIC ACTION

Tear stain cleaner.

INGREDIENTS

Purified water, Propylene Glycol, Cocamidopropyl Betaina, Allantoina, Isothiazolinones.

MODE OF APPLICATION

1. Moisten a cotton and apply it gently over the affected (stained) area, avoiding contact with the eye surface.
2. Then dry with a dry cotton.
3. The areas very stained will require treatment on a daily basis, until the stains have been removed.
4. When the stains have disappeared, treat regularly once a week to prevent the recurrence.

WARNINGS

- Mantener fuera del alcance de los niños
- En caso de contacto accidental con los ojos, lavar con abundante agua.

OBSERVATIONS

Do not have irritant substances.

CONSERVATION

Store in a cool, dry place, protected from light, below 30 ºC.

PRESENTATION

50 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

LAXDRAG® ORAL PASTE

PASTA ORAL.

TRATAMIENTO Y PREVENCIÓN DE LAS BOLAS DE PELO Y CONSTIPACIÓN EN GENERAL.

Technical Specification

SPECIES

Cats and dogs.

- Prevent constipation problems.
- Delicious salmon flavor.
- Prevent obstructive gastrointestinal diseases.

DOSAGE FORM

Oral Paste.

THERAPEUTIC ACTION

Prevent constipation problems.

INGREDIENTS

Malt extract, petrolatum, glycerin, salmon flavor, acacia and vitamin B1.

PROPERTIES

When cats lick themselves, their tongues remove hairs from fur, which are ingested and accumulated in the gastrointestinal tract of animal. This material, practically indigestible, becomes “hairballs”, which may interfere in the digestion and cause problems with its elimination. The symptoms of your pet with “Hairballs” are:

- constipation
- difficult bowel motion
- dry cough and vomiting after meals.

The use of LAX DRAG®, and the regular brushing of your pet can eliminate these problems.

ROUTE OD ADMINISTRATION AND DOSAGE

a. To eliminate “Hairballs”:

- Apply daily 2 to 3 cm of LAX DRAG® to your adult cat until the symptoms disappear.
- Administer it between meals.
- For applying it, put the quantity indicated in your finger or directly on one of the hands of your pet, from where he will easily lick it.

b. To prevent “Hairballs”:

- Give your adult cat 2 to 3 cm of LAX DRAG® once or twice a week, and also brush it regularly.
- Fore kittens over 4 weeks old: 1 to ½ cm of LAX DRAG®, once or twice a week.
- In dogs suffering occasional constipation, administer 2 to 3 cm of LAX DRAG®, once or twice a week.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

Salmon flavor.

CONSERVATION

Store in a fresh, dry place, out of direct light, at no more than 30 °C.

PRESENTATION

90 g

PREPARED BY

Drag Pharma Laboratory.

LIDOCALM® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANESTÉSICO LOCAL.

Technical Specification

SPECIES

Horses, cattle, sheep, goats, pigs, dogs and cats.

It is recommended as local anesthetic by infiltration or neural blockade in interventions such as: castration, suture of wounds, ophthalmic intervention, vulvoplasty or any other intervention requiring temporary suspension of pain sensation. It is recommended for perineural blockade for the diagnosis of lameness; as paravertebral or medulla anesthetic, for ruminitomies or cesarean sections.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Local anesthetic.

COMPOSITION

Each 1 mL of solution contains:
Lidocaine hydrochloride ..... 20.0 mg
Excipients q.s.p.....1 mL

INDICATIONS

- Do not use in animals with known hypersensitivity to local anesthetics of the amide class.
- Do not administer in pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Subcutaneous, intramuscular and epidural administration, in all species.

Dose of the active principle:	**Local Anesthesia**	**Epidural Anesthesia**
Specie		
Horses	Maximum 4 mg/Kg	0,2 mg/Kg
Bovines	Maximum 4 mg/Kg	0,2 mg/Kg
Sheep and goats	Maximum 4 mg/Kg	4,4 mg/Kg
Pigs	Maximum 4 mg/Kg	4,4 mg/Kg (máx 400 mg)
Dogs	Maximum 8 mg/Kg	4,4 mg/Kg
Cats	Maximum 4 mg/Kg	4,4 mg/Kg

Product dose:	**Local Anesthesia**	**Epidural Anesthesia**
Specie		
Horses	Maximum 20 mL/100 Kg	1 mL/100 Kg
Bovines	Maximum 20 mL/100 Kg	1 mL/100 Kg
Sheep and goats	Maximum 4 mL/20 Kg	1 mL/4,5 Kg
Pig	Maximum 4 mL/20 Kg	1 mL/4,5 Kg (maximum 20 mL)
Dogs	Maximum 4 m /10 Kg	1 mL/4,5 Kg
Cats	Maximum 0,2 mL/Kg	1 mL/4,5 Kg

CONTRAINDICATIONS

- Do not use in animals with known hypersensitivity to local anesthetics of the amide class.
- Do not administer in pregnant or lactating females.

PRECAUTIONS

Special warnings and precautions for use:

- The dose required will depend on the extent of the region to be anesthetized, the type and magnitude of the intervention, and the species.
- Use with caution in animals with liver or heart dysfunction.
- Avoid an overdose of the product, it can cause depression of neuromuscular and / or cardiac functions.
- Cats tend to be more sensitive to the effects of the drug on the central nervous system, administer with caution.
- Lidocaine toxicity in dogs is enhanced by Phenobarbital.
- Accidental intravenous injection should be avoided.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR USE

Special warnings and precautions for use:

- The dose required will depend on the extent of the region to be anesthetized, the type and magnitude of the intervention, and the species.
- Use with caution in animals with liver or heart dysfunction.
- Avoid an overdose of the product, it can cause depression of neuromuscular and / or cardiac functions.
- Cats tend to be more sensitive to the effects of the drug on the central nervous system, administer with caution.
- Lidocaine toxicity in dogs is enhanced by Phenobarbital.
- Accidental intravenous injection should be avoided.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Avoid contact with the skin. Wash hands with soap and water after handling the product.
- In case of accidental self-injection, seek medical attention immediately and show the internal leaflet to the doctor.
- People with known hypersensitivity to Lidocaine Hydrochloride should avoid contact with the veterinary drug.

GUARD PERIOD

Meat: 28 days.
Milk: 7 days.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Dispose of unused product remains in its original container, well closed. Do not dispose of containers with product remains on the ground or water courses. Dispose of the waste of this product with care together with household waste.

CONSERVATION

Store between 15 and 30 ° C, protected from light. Use immediately once opened and discard the excess product.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Ampule vial with 50 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No. 1865

LOMBRIMIC® ORAL SUSPENSION

SUSPENSIÓN ORAL.

ANTIPARASITARIO INTERNO DE AMPLIO ESPECTRO.

Technical Specification

SPECIES

Bovine, sheep, goats, pigs and horses.

It is indicated for the treatment of mature and immature forms of gastrointestinal and pulmonary parasites and tapeworms, including ovicidal action.

DOSAGE FORM

Oral suspension.

THERAPEUTIC ACTION

Broad spectrum internal antiparasitic.

COMPOSITION

Each 100 ml contains:
Fenbendazole.....10 g
Excipients q.s.p.....100 mL

INDICATIONS

Do not administer in animals with known hypersensitivity to Fenbendazole.

ACTION SPECTRUM

Bovine, sheep and goats: *Haemonchus spp., Cooperia spp., Ostertagia spp., Trichostrongylus spp., Oesophagostomum spp., Strongyloides spp., Trichuris spp., Bunostomum spp., Dictyocaulus spp., Nematodirus spp., Chabertia ovina, Moniezia spp.*

Horses: Large and small strongyles, *Parascaris equorum, Trichonema spp., Trichostrongylus axei, Strongyloides westeri.*

Pigs: *Oesophagostomum spp., Metastrongylus spp., Trichuris suis, Ascaris suum, Strongyloides ransomi.*

ROUTE OD ADMINISTRATION AND DOSAGE

Administer orally through a dosing gun or syringe.

Dose:

- **Cattle, sheep, goats, pigs:** 5 mL / 100 Kg of weight (Equivalent to 5 mg of Fenbendazole / Kg of weight). Single dose.
- **Horses:** 7.5 mL / 100 Kg of weight in a single dose (Equivalent to 7.5 mg of Fenbendazole / Kg of weight). Single dose.

CONTRAINDICATIONS

Do not administer in animals with known hypersensitivity to Fenbendazole.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not administer in animals with known hypersensitivity to Fenbendazole. Wear gloves when administering the product.
- Wash your hands after handling the product.
- Do not eat, drink or smoke while handling the product.
- Do not handle by people who are hypersensitive to Fenbendazole.
- Avoid contact with skin, eyes, or mouth.
- In case of accidental ingestion, go immediately to a medical center and show the product label.
- In case of contact with skin, eyes or mouth, wash immediately with plenty of water.
- In case of skin irritation, go immediately to a medical center and show the product label.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

- **Cattle:** Meat: 14 days. Milk: 7 days
- **Horses:** Do not administer in animals whose meat is intended for human consumption
- **Sheep and goats:** Do not administer in animals whose milk is intended for human consumption.

OBSERVATIONS

Shake vigorously before use.

Special precautions for unused product or waste material:

- Empty containers can be disposed of as household waste, without any special precautions.
- Do not dispose of containers with product remains on the ground or water courses.
- For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30°C, protected from light.
Once the container is opened, use within 12 weeks. Discard unused product after that time frame.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Bottle with 100 ml, 500 ml, 1 L and 3 L

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 0624
- Dominican Republic: Reg. No. 6516

MAMISTOP® CAT - ORAL POWDER

POLVO ORAL.

SUSTITUTO LÁCTEO FELINO

Technical Specification

SPECIES

Cats.

Complete and balanced formula for kittens and cats subjected to stressful conditions.

DOSAGE FORM

Oral powder.

THERAPEUTIC ACTION

Feline dairy substitute

COMPOSITION

GUARANTEED ANALYSIS (as powder):
Crude protein, min 40%
Crude fat min 21%
Crude fiber 0%
Humidity, max 5%
Taurine..... 0.1%
Calcium (min - max) ..... 0.2 - 0.7%
Phosphorus (min - max) ..... 0.4 - 1%

INGREDIENTS

Powdered whole cow's milk, Powdered dehydrated egg, Sodium Caseinate, Powdered skimmed cow's milk, Lactose, Taurine, Minerals (Zinc (as Sulfate), Manganese (as Sulfate), Iron (as Sulfate), Antioxidant agents and preservatives.

PROPERTIES

MamiStop® Cats is a complete and balanced formula that provides a nutritional profile similar to that of feline breast milk. Indicated as a food source for orphaned, rejected, growing kittens or as a nutritional supplement in adult cats subjected to stressful situations (pregnant, lactating, convalescent, show cats) that require the contribution of highly digestible nutrients.

USE INSTRUCTIONS

Method of preparation and dosage:

- **Newborn and growing kittens:** Dissolve 1 tablespoon of powder in 2 tablespoons of warm boiled water. Administer 20 mL for every 100 g of weight per day, divided into 4 doses; or according to Veterinary Doctor indication. After administering the product, it is recommended to stimulate the perineum with a soft, moist and warm cloth to stimulate urination and defecation.
- **Pregnant or lactating cats:** Dissolve 2 tablespoons of powder in 4 tablespoons of warm boiled water. Administer 60 mL for every 2 kg of weight per day directly orally, or mixed with food for up to two weeks after delivery.
- **Show or convalescent cats:** Dissolve 1 tablespoon of powder in 2 tablespoons of warm boiled water. Administer 30 mL for every 2 Kg of weight per day directly orally, or mixed with food.

(*) 1 tablespoon = 15 g of powder / 15 mL of water.

Follow the preparation instructions and give only the recommended amount to avoid digestive discomfort.

ROUTE OD ADMINISTRATION AND DOSAGE

The recommended dose of the reconstituted product is 4 teaspoons (20 mL) daily per 100 g of weight, or as indicated by the Veterinarian. In kittens of 0 to 2 weeks of age, it is suggested to feed 4 times a day, dividing the total volume in 4 doses; in older kittens, feed 3 times a day.

Age (weeks)	Mean weight (g) *	Volumen/day	Number of doses/day
1	115	20 mL	4
2	200	40 mL	4
3	280	50 mL	3
4	375	65 mL	3
5	480	80 mL	3

* Kittens increase their weight in 50-100 g per week.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

VETERINARY USE
EXCLUSIVE USE IN ANIMAL FEED
FORBIDDEN USE IN THE FEEDING OF RUMIANTS.

CONSERVATION

Keep in a fresh, dry place, out of the direct light, at no more than 30°C.

CONDITION OF SALE

Free sales (OTC)

PRESENTATION

Pot with 100 g

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. LENAA N°: RM 03-008N
Costa Rica: Lic. DAA-MAG 579-005
Bolivia: Reg. SENASAG PUV-A-N° 005513/13
Perú: Reg. SENASA A.001.002.I.05863

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Costa Rica by:
Proventas de Cartago S.R.L. 100 meters east Hogares Crea, San Blas. Carthage.
Tel: 2591 4624 - Fax: 2591 5339

Imported and distributed in Bolivia by:
AGROGUARANI SRL.
TEL: + (591) 314-1401
Santa Cruz de la Sierra, Bolivia

MAMISTOP® DOG- ORAL POWDER

POLVO ORAL.

SUSTITUTO LÁCTEO CANINO

Technical Specification

SPECIES

Dogs.

Complete and balanced formula for puppies and dogs subjected to stressful conditions.

DOSAGE FORM

Oral powder.

THERAPEUTIC ACTION

Canine dairy substitute

COMPOSITION

GUARANTEED ANALYSIS (as powder):
Crude protein (min) 20%
Crude fat (min) 20%
Crude fiber (max) 0%
Humidity (max) 5%
Calcium (min - max) .....1% - 1.5%
Phosphorus (min - max) .... 0.8% - 1.2%

INGREDIENTS

Whole cow's milk powder (26% fat), sugar, dehydrated egg powder, Calcium Hydrogen Phosphate, authorized sweetening agents and preservatives.

PROPERTIES

MamiStop® Dogs is a complete and balanced formula that provides a nutritional profile similar to that of canine breast milk. Indicated as a food source for orphaned puppies, large litters, weaning period, early weaning, inability of the female to feed her litter (mastitis, cesarean section, etc.) or as a nutritional supplement in adult dogs subjected to stressful situations (pregnant , infants, convalescents, malnutrition, show dogs) that require the contribution of highly digestible nutrients.

USE INSTRUCTIONS

- **For puppies:** Dissolve 15 g of powder (*) in 50 mL of warm boiled water. Shake until completely dissolved and administer on demand (ad libitum) in several doses a day or as indicated by a Veterinary Doctor. After administering the product, it is recommended to stimulate the perineum with a soft, moist and warm cloth to stimulate urination and defecation.
- **Adult dogs:** Dissolve 30 g of powder (*) in every 100 mL of warm boiled water. Administer 100 mL for every 10 Kg of weight directly orally or mixed with food.

(*) 15 g of powder = 1 tablespoon.

Follow the preparation instructions and administer only the recommended amount to avoid digestive discomfort.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

EXCLUSIVE USE IN ANIMAL FEED
FORBIDDEN USE IN THE FEEDING OF RUMIANTS

CONSERVATION

Keep in a cool, dry place and protected from light, at no more than 30ºC.

CONDITION OF SALE

Free Sales (OTC)

PRESENTATION

Pot with 125 g and 250 g

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. LENAA N°: RM 03-008N
Panamá: RF-8468-19
Costa Rica: Lic. DAA-MAG 579-004
Bolivia: Reg. SENASAG PUV-A-N° 005514/13
El Salvador: AL2007101602
Perú: Reg. SENASA A.36.15.I.0039

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Costa Rica by:
Proventas de Cartago S.R.L. 100 meters east Hogares Crea, San Blas. Carthage.
Tel: 2591 4624 Fax: 2591 5339

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

Dist. In El Salvador by Rafael Alfredo Alfaro Castillo.

Imported and distributed in Peru by Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

MASTERFLY® TOPICAL SOLUTION

SOLUCIÓN TÓPICA

ANTIPARASITARIO EXTERNO - MOSQUICIDA POUR ON.

Technical Specification

SPECIES

Bovine.

MASTERFLY is an external antiparasitic indicated for the control and elimination of horn flies (Haematobia irritans) in cattle. Field studies demonstrated a maximum efficacy of 35 days in the control of horn flies, except in conditions in which the animals are exposed to extreme rain conditions in which the efficacy decreases to 21 days.

DOSAGE FORM

Topical solution.

THERAPEUTIC ACTION

External anti-parasite - Fly killer pour on.

COMPOSITION

Each 100 mL of solution contains:
Permethrin.....10 g
Cypermethrin.....7 g
Piperonyl Butoxide.....20 g
Excipients q.s.p.....100 mL

INDICATIONS

- Do not administer in dairy cattle.
- Do not administer during pregnancy and lactation.

EFFECTIVENESS

ROUTE OD ADMINISTRATION AND DOSAGE

Route of administration: Topical

The recommended dose to be administered to cattle is 15 mL for animals up to 250 Kg of body weight and 20 mL for animals more than 250 Kg of body weight, single dose. It is recommended to treat the whole animal mass to better control the horn fly.

It is recommended applying the product with a dispensing gun in a dorsal longitudinal line against the grain, from the base tail (rump) toward the withers, to facilitate the contact of the product and the animal skin.

CONTRAINDICATIONS

- Do not administer in dairy cattle.
- Do not administer during pregnancy and lactation.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- The product is toxic if ingested and harmful in case of contact with the skin and eyes (Irritant)
- The operator should wear gloves and protective gears to handle this product.
- Avoid contact with the skin, eyes and clothing.
- Do not smoke, eat or drink during product application.
- Never transfer the product to another container for storage.
- Wash hands with plenty soap and water after handling the product and before eating, drinking or smoking.
- Wash contaminated clothes before reuse.
- In case of eye contact, immediately flush eyes with water for at least 15 minutes.

WARNINGS

- Mantener fuera del alcance de los niños.
- Mantener alejado de los alimentos.
- Mantener en su envase original y con su etiqueta visible.

GUARD PERIOD

Meat: 14 days.

Milk: Do not administer to milk producing cattle.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:

- Once opened, discard the unused product in its container.
- Do not discard the packaging and/or product residues in rivers, lakes or streams of water.
- Do not throw away empty containers or container with product residues with the household waste.
- Do not reuse container.
- Dispose this product with companies that provide special waste removal services.

CONSERVATION

Store at a temperature between 2 ° C and 30 ° C, protected from light. Once the container is opened, the product is stable between 15 ° C and 30 ° C for 12 weeks.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

1 L and 3 L container

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG Nº 2199
Bolivia: Reg. SENASAG PUV-F N° 007249/16
Perú: Registro SENASA F.87.33.I.0183

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:
AGROGUARANI SRL
TEL: + (591) 314-1401
Santa Cruz de la Sierra, Bolivia

MATIHORSE® TOPICAL GEL

GEL TÓPICO.

AYUDA EN EL PROCESO DE RESTAURACIÓN DE LA PIEL.

Technical Specification

SPECIES

Horses.

Its use is helpful in restoration processes of the skin.

DOSAGE FORM

Topical gel.

THERAPEUTIC ACTION

Help in the recovery process of the skin.

COMPOSITION

Matico standardized extract.

Excipients q.s.p. 100 g

PROPERTIES

Matihorse® is a cream made from natural Matico (*Buddleja globosa*) extract, which has been specially formulated for use in horses and companionship animals.

ROUTE OD ADMINISTRATION AND DOSAGE

Apply twice a day after cleaning the affected area.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a cool and dry place, at room temperature between 15º and 30ºC.

PRESENTATION

Pomander with 100 g and container with 500 g

PREPARED BY

Drag Pharma Laboratory.

MATIPET® CREAM

CREMA.

CREMA NATURAL DE MATICO, CALÉNDULA Y ÁRNICA.

Technical Specification

SPECIES

Dogs and cats.

MATIPET® is a cream manufactured from natural extracts of Matico (*Buddleja globosa*), Marigold (*Caléndula officinalis*) and Arnica (*Arnica montana*), which has been specially formulated to be used in dogs and cats.

DOSAGE FORM

Cream.

THERAPEUTIC ACTION

Natural Cream of Matico, Marigold and Arnica.

COMPOSITION

Each 100 g of cream contains:

Matico extract .....10 g

Marigold extract .....5 g

Arnica extract .....5 g

Vitamin E acetate .....4 g

Zinc oxide .....3 g

Excipients q.s.p. .....100 g

MODE OF APPLICATION

1. Clean the wound or affected area with warm water, removing crusty secretions and cell debris.
2. Apply the cream covering completely the affected zone, let act and do not remove it.
3. Apply the product 2 to 3 times daily, as necessary.

ROUTE OD ADMINISTRATION AND DOSAGE

Topical cream.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a fresh, dry place, out of direct light, at no more than 30 °C.

PRESENTATION

50 g

PREPARED BY

Drag Pharma Laboratory.

MEBERMIC® ORAL TABLETS

COMPRIMIDO ORAL.

ANTIPARASITARIO INTERNO.

Technical Specification

SPECIES

Dogs and cats.

Treatment of all internal parasites (Nematodes and Cestodes) of dogs and cats (except *Trichuris vulpis*).

DOSAGE FORM

Oral tablets.

THERAPEUTIC ACTION

Internal antiparasitic

COMPOSITION

Each tablet contains:
Praziquantel.....50 mg
Mebendazole.....220 mg
Excipients q.s.p.....1 tablet

INDICATIONS

- Do not use in dogs less than 4 weeks old or in cats less than 6 weeks old.
- Do not use in animals with impaired hepatic, renal or cardiac function.
- Do not administer during pregnancy or while breastfeeding.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration.

Dose:
1 tablet / 10 Kg of body weight, in a single dose (equivalent to 5 mg / Kg of Praziquantel and 22 mg / Kg of Mebendazole).

DRUG INTERACTIONS

Do not use concomitantly with other pharmaceutical products.

CONTRAINDICATIONS

- Do not use in dogs less than 4 weeks old or in cats less than 6 weeks old.
- Do not use in animals with impaired hepatic, renal or cardiac function.
- Do not administer during pregnancy or while breastfeeding.

WARNINGS

Mantener fuera del alcance de los niños.

SIDE EFFECTS

In some animals, hypersalivation, nausea, vomiting, depression, loose stools or diarrhea may occur temporarily.

CONSERVATION

Keep in a cool and dry place, at room temperature between 15 and 30° C

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Box with 50 tablets.

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG. N°: 0451
Perú: Reg. SENASA F.08.21.I.1125

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Peru by Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

MELOXIVET® ORAL SOLUTION

SOLUCIÓN ORAL.

ANALGÉSICO Y ANTIINFLAMATORIO.

Technical Specification

SPECIES

Dogs

Meloxivet®, Meloxicam 1 mg / mL, Oral solution, is indicated for the control of pain and inflammation associated with osteoarthritis symptoms in dogs.

DOSAGE FORM

Oral solution

THERAPEUTIC ACTION

Analgesic and Anti-inflammatory.

COMPOSITION

Each mL of oral solution contains:
Meloxicam ..... 1 mg
Excipients q.s.p ..... 1 mL

TECHNICAL CHARACTERISTICS

INDICATIONS

- Meloxicam is contraindicated in dogs with known hypersensitivity to Meloxicam, Piroxicam or other NSAIDs.
- Do not administer to dogs younger than 6 months or weighing less than 6 kilos.
- Meloxicam should not be used in dogs with ulceration or active gastrointestinal bleeding.
- Do not use in patients with liver, heart or kidney failure and bleeding disorders.
- Do not administer to pregnant or lactating females or breeding animals.

MODE OF APPLICATION

The solution must be administered using the dosing syringe included in the package, according to the weight of each animal. Administer mixed with a portion of food or directly into the dog's muzzle.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration.

Administer an initial dose of 0.2 mL for each kilo of weight on the first day of treatment, followed by a maintenance dose of 0.1 mL for each kilo of weight every 24 hours. These doses are equivalent to administering an initial dose of the active ingredient of 0.2 mg / kg of weight on the first day of treatment, followed by a maintenance dose of 0.1 mg / kg of weight every 24 hours.

The duration of treatment will depend on the condition to be treated. For acute pain, the duration of treatment can last up to 15 days depending on the degree of pain and inflammation. For chronic pain, the duration of treatment can be prolonged from 21 to 28 days. Both acute and chronic treatment should be discontinued after 10 days if there is no apparent improvement or if vomiting or occult blood occurs in the stool.

DRUG INTERACTIONS

- Nonsteroidal anti-inflammatory drugs (NSAIDs) may reduce the effects of angiotensin-converting enzyme inhibitors (eg Enalapril, Benazepril) on blood pressure.
- Meloxicam interacts with anticoagulants like Heparin or Warfarin and may increase the possibility of bleeding.
- Its administration with corticosteroids or other NSAIDs could increase the risk of gastrointestinal toxicity (ulceration, bleeding, vomiting or diarrhea).
- NSAIDs may reduce the diuretic effects of Furosemide.
- NSAIDs may increase the risk of nephrotoxicity, when used in conjunction with nephrotoxic or diuretic drugs, such as Furosemide and aminoglycosides.

CONTRAINDICATIONS

- Meloxicam is contraindicated in dogs with known hypersensitivity to Meloxicam, Piroxicam or other NSAIDs.
- Do not administer to dogs younger than 6 months or weighing less than 6 kilos.
- Meloxicam should not be used in dogs with ulceration or active gastrointestinal bleeding.
- Do not use in patients with liver, heart or kidney failure and bleeding disorders.
- Do not administer to pregnant or lactating females or breeding animals.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- En caso de contacto con la piel se recomienda lavar las manos con jabón y abundante agua.
- En el caso de contacto con los ojos, lávese con abundante agua por 15 minutos. Si se desarrolla irritación y esta persiste, consultar al médico.
- En caso de ingestión accidental, no inducir el vómito. Lave la boca con abundante agua. En caso de desarrollar irritación gástrica, obtener ayuda médica.

WARNINGS

Advertencias y precauciones especiales de uso:

- Se debe tener especial cuidado en animales deshidratados, hipovolémicos e hipotensos ya que existe un mayor riesgo potencial de desarrollar toxicidad renal.
- El tratamiento debe ser suspendido en el caso de no observarse la respuesta clínica esperada.
- Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

- Adverse effects reported in dogs treated with Meloxicam are similar to those observed with other NSAIDs and include gastrointestinal effects (vomiting, loss of appetite, loose stools, diarrhea, melena, gastrointestinal ulcerations, hematemesis), liver (elevated liver enzymes, jaundice), dermatological (pruritus, urticaria, dermatitis), kidney (azotemia, creatinine and blood urea elevation, renal failure), central nervous system and behavior (ataxia, personality disorders, seizures, drowsiness, hyperactivity, depression, tremors and lethargy in puppies) and hematological (alteration in coagulation times, immunomediated hemolytic anemia, immunomediated thrombocytopenia).
- The unwanted effects generally occur after 7 days of treatment and in most cases are transient in nature and disappear once treatment is complete. In the case of observing unwanted effects and / or adverse reactions, treatment should be suspended.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
All unused medicine or the residues derived from it, must be disposed of in an environmentally safe way. Do not dispose of empty containers or containers with product residues in water courses. Do not throw away empty containers or containers with product residues along with household waste. Contact the manufacturing laboratory to receive instructions regarding the correct disposal.

CONSERVATION

Store at a temperature between 15 and 30 ° C, protected from light. Once opened, use the product within 30 days. Discard the unused product after that period of time.

CONDITION OF SALE

Sale with a Veterinary Medical prescription.

PRESENTATION

Bottle with 60 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG Nº 2415
Perú: Registro. SENASA F.99.32.I.0130

AVAILABLE FOR SALE

Imported and Distributed in Peru by:
Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

LEVEL

1

METAMIZOL SÓDICO (DIPIRONA) - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANALGÉSICO - ANTIINFLAMATORIO - ANTIPIRÉTICO.

Technical Specification

SPECIES

Bovine, horses, pigs, sheep, goats and cats.

Analgesic, anti-inflammatory and antipyretic.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Analgesic – Anti-inflammatory – Antipyretic.

COMPOSITION

Each 100 mL contains.

Metamizole Sodium..... 50 g

Excipients q.s.p.....100 mL

INDICATIONS

- Do not use in conjunction with chlorpromazine, barbiturates and/or Phenylbutazone.
- Do not use in animals with a history of blood dyscrasias.

ROUTE OD ADMINISTRATION AND DOSAGE

Slow intramuscular or intravenous administration.

- Bovine and horses: 20 mL/400 Kg
- Foals: 10 mL/150 Kg
- Pigs: 5 mL/100 Kg
- Dogs: 0.4 mL/10 Kg
- Cats. 0.2 mL/5 Kg

Repeat the dose 2 to 3 times a day.

CONTRAINDICATIONS

- Do not use in conjunction with chlorpromazine, barbiturates and/or Phenylbutazone.
- Do not use in animals with a history of blood dyscrasias.

PRECAUTIONS

- The application of the product for long periods of time, requires periodic evaluations of the patient.
- Metamizole overdose may cause convulsive events.
- Use with caution in aged animals and/or animals with cardiovascular diseases.
- Metamizole Sodium may tend to increase the likelihood of bleeding, due to suppression in the prothrombin formation.

SPECIAL PRECAUTIONS FOR USE

- The application of the product for long periods of time, requires periodic evaluations of the patient.
- Metamizole overdose may cause convulsive events.
- Use with caution in aged animals and/or animals with cardiovascular diseases.
- Metamizole Sodium may tend to increase the likelihood of bleeding, due to suppression in the prothrombin formation.

WARNINGS

- Mantener fuera del alcance de los niños
- En animales de competencia, usar solo 5 días antes de la competencia.
- La administración vías subcutánea puede causar irritación en el sitio de inyección.
- La aplicación del producto por tiempos prolongados, requiere evaluación periódica del paciente.
- La sobredosis de Metamizol, puede provocar eventos convulsivos.

GUARD PERIOD

Meat: Bovine, sheep and pigs: 35 days.

Horses: Do not administer to horses which meat is intended for human consumption.

Milk: Do not administer to animals which milk is intended for human consumption.

CONSERVATION

Store between 15 and 30 °C, protected from light. Once the container is opened, use within 28 days. Discard unused product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Ampule vial with 50 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

- Chile: Reg. SAG No. 0528
- Panama: Reg. No. RF-3901-07

MINERICE® TOPICAL GEL

GEL TÓPICO.

AYUDA PARA LAS MOLESTIAS MUSCULARES Y ARTICULARES.

Technical Specification

SPECIES

Horses.

Help for muscle and joint discomfort in horses under great physical demands.

DOSAGE FORM

Topical gel.

THERAPEUTIC ACTION

Help for muscle and joint discomfort.

COMPOSITION

a) Active: Peppermint-camphor, 2-Camphanone, Wintergreen essential oils.

b) Other: Copper, Isopropyl alcohol, Magnesium, Sodium and special carriers.

PROPERTIES

Minerice is a pleasantly refreshing gel, non-greasy.

INDICATIONS

- Do not apply other products in the same area where Minerice® will be apply.
- Do not roughly scrub or massage.
- Do not apply over open wounds or damage skin,
- Do not put bandages too tight.

ROUTE OD ADMINISTRATION AND DOSAGE

Topical use.

Directions:

- The surface area to be treated should be clean and free of previously applied products.
- Apply Minerice® with the hand or a wet towel, covering completely the affected area. To get a good result, carefully massage to get a good contact with the animal skin. It is convenient to put bandages on the treated extremity or cushioning with resting bandages.

How often to apply?

It is recommended to apply at least 2 times a day, before and after physical activity.

When to apply?

- Training horses: Apply before and after sport work.
- After athletic competition or major effort demanding work: Apply in helping to relieve discomfort as consequence of high demand competitions such jumping competitions, training, polo, racing, complete race, endurance, long trips, rodeos, etc.
- Transporting horses: Apply Minerice® and place transport protections. The horse will feel pleased and will arrive in better conditions for the competition.
- As refreshing bath: It can be used as refreshing and revitalizing bath if is used as follows: Dilute 10 to 15 g
- (1 tablespoon) of Minerice®, into a container with 400 ml (2 cups) of water. Mix vigorously to form a soft and even cream. Then mix this just formed cream with approximately 5 L of water. Apply this dilution to the animal body with a sponge and then rinse, pat the horse coat and he will feel physically great and emotionally pleased.

CONTRAINDICATIONS

- Do not apply other products in the same area where Minerice® will be apply.
- Do not roughly scrub or massage.
- Do not apply over open wounds or damage skin,
- Do not put bandages too tight.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° and 30°C.

PRESENTATION

Container with 500 g

PREPARED BY

Drag Pharma Laboratory.

MIXANTIP® PLUS - CREAM

CREMA.

**ANTIBIÓTICO, ANTIALÉRGICO, ANTIINFLAMATORIO,
ANTIMICÓTICO, ANTIPRURIGINOSO, ANESTÉSICO Y CICATRIZANTE.**

Technical Specification

SPECIES

Dogs and cats.

- Help in the treatment of bacterial and fungal infections, painful and diffuse skin inflammations and itching in companion animals.
- Dermatitis, eczema, pyoderma and abrasions.
- Cosmetic surgeries (tail docking, ear cropping).

DOSAGE FORM

Cream.

THERAPEUTIC ACTION

Antibiotic, Anti-inflammatory, Antiallergy, Antifungal Antipruritic, Anesthetic and healing.

COMPOSITION

Each 100 g contains:
Prednisolone Acetate.....0.1117 g
(Equivalent to 0.1 g of Prednisolone base)
Neomycin Sulfate.....500.000 UI
Clotrimazole.....1.00 g
Precipitated Sulfur.....0.50 g
Zinc Oxide.....3.00 g
Benzocaine.....1.00 g
Excipients q.s.p.....100.0 g

ROUTE OD ADMINISTRATION AND DOSAGE

Topically.

Clean and shave the affected area, gently spread a sufficient amount of cream on the area 2 to 3 times a day, for 5 to 7 days, or until complete recovery of the wound.

WARNINGS

- No usar en animales con hipersensibilidad a algunos de sus componentes.
- Administrar con precaución en animales en gestación.
- Mantener fuera del alcance de los niños.

CONSERVATION

Store in a cool and dry place between entre 15° and 30°C, protected from sunlight.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Pot with 15 g and 50 g

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No. 1597-B

Perú: Reg. SENASA F.102.039.I.00004

MOSKIMIC® TOPICAL SOLUTION

SOLUCIÓN TÓPICA.

ANTIPARASITARIO *POUR ON.*

Technical Specification

SPECIES

Bovines

MOSKIMIC® Pour On, is an external antiparasitic, which can be administered to cattle of any age. It has a wide spectrum of activity, repels, eliminates and controls flying insects (horn flies, flies in general, horseflies, mosquitoes, etc.). It is also recommended for its lousy effect.

DOSAGE FORM

Topical Solution

THERAPEUTIC ACTION

Antiparasitic *pour on*

COMPOSITION

Each 100 mL contains:
Permethrin ..... 10 g
Piperonyl Butoxide ..... 10 g
Excipients q.s.p ..... 100 mL

ROUTE OD ADMINISTRATION AND DOSAGE

Topical application.

- Against flies: Apply 10 mL per animal.
- Against lice: Apply 10 mL per animal up to 100 kg in weight. Apply 20 mL per animal between 100 to 300 Kg of weight.

With a dosing gun or syringe, apply in a dorsal longitudinal line from the base of the tail (rump) to the region of the withers. Repeat the treatment at 4 weeks.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Overexposure and / or in subjects susceptible to the product, may cause side effects (allergies, skin rush, others).
- When applying the product, wear gloves.
- In case of accidental contact, wash the affected area with plenty of soap and water.
- In case of poisoning, consult a doctor.

WARNINGS

- Mantener fuera del alcance de los niños.
- No aplicar próximo a fuentes de calor.
- No desechar el envase en ríos.
- Evitar el contacto con los ojos.
- Compatible con otros tratamientos como vacunaciones y desparasitaciones.

GUARD PERIOD

Meat: 0 days.

Milk: 2 days.

CONSERVATION

Store in a cool and dry place, at room temperature between 15 ° and 30 ° C.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

Container with 100 mL, 250 mL, 1 Liter and 3 Liters.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG Nº 625

NANORMEN® ORAL SUSPENSION

SUSPENSIÓN ORAL.

ANTIPARASITARIO NEMATICIDA

Technical Specification

SPECIES

Dogs and cats.

Nanormen® Oral Suspension, is indicated for the treatment and control of gastrointestinal nematodes in dogs and cats of any age.
Nanormen® is effective against *Toxocara canis, Toxocara cati, Toxascaris leonina, Ancylostoma caninum, Ancylostoma tubaeforme and Uncinaria stenocephala.*

DOSAGE FORM

Oral suspension

THERAPEUTIC ACTION

Nematicidal dewormer

COMPOSITION

Each mL contains:
Pirantel Pamoate ..... 144.13 mg
(Equivalent to 50 mg of Pirantel base)
Excipients q.s.p ..... 1 mL

INDICATIONS

- Do not administer to severely weakened animals.
- Do not administer to pregnant or lactating females.
- Do not administer to hypersensitive animals to Pirantel Pamoato.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration way: Oral.

Dose of the active substance:

- Dogs: 5 mg / Kg of weight.
- Cats: 20 mg / Kg of weight.

Product dosage:

- **Dogs: administer 2 drops per Kg of weight or 1 mL for every 10 Kg of weight in a single dose.**
- **Cats: Administer 8 drops per Kg of weight or 2 mL for every 5 Kg of weight in a single dose.**

DRUG INTERACTIONS

Pirantel can interact with medications containing Piperazine, Levamisole, or other Nicotine-like cholinergic drugs.

CONTRAINDICATIONS

- Do not administer to severely weakened animals.
- Do not administer to pregnant or lactating females.
- Do not administer to hypersensitive animals to Pirantel Pamoato.

PRECAUTIONS

Special warnings and precautions for use:

- Shake before using.
- Keep out of the reach of children.
- Nanormen® is recommended to treat and control parasite reinfestation in puppies and kittens, for which animals can be treated at 2, 4, 6, 8 and 10 weeks of life.
- Animals exposed to highly contaminated environments, it is recommended to administer Nanormen® every 2 or 3 months to control reinfestation by *Toxocara canis.*

SPECIAL PRECAUTIONS FOR USE

Special warnings and precautions for use:

- Shake before using.
- Keep out of the reach of children.
- Nanormen® is recommended to treat and control parasite reinfestation in puppies and kittens, for which animals can be treated at 2, 4, 6, 8 and 10 weeks of life.
- Animals exposed to highly contaminated environments, it is recommended to administer Nanormen® every 2 or 3 months to control reinfestation by *Toxocara canis.*

SPECIAL PRECAUTIONS FOR THE OPERATOR

Do not handle by people hypersensitive to Pirantel Pamoato

WARNINGS

SIDE EFFECTS

Mild and transient side effects may occasionally occur, including gastrointestinal signs such as nausea, vomiting, anorexia, abdominal pain, diarrhea, erythema of the skin, and increased liver enzyme values.

ADVERSE EFFECTS

Mild and transient side effects may occasionally occur, including gastrointestinal signs such as nausea, vomiting, anorexia, abdominal pain, diarrhea, erythema of the skin, and increased liver enzyme values.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
Empty containers can be discarded as household waste, without any special precautions. Do not dispose of containers with product residues on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 2 ° and 30 ° C, protected from light. Once opened, use within 12 weeks and discard the excess product.

CONDITION OF SALE

sale with Veterinary Medical prescription

PRESENTATION

20 mL bottle

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 178
Costa Rica: Reg. Nº MAG CL4-42-10-5936

NANORMEN® PLUS - ORAL TABLET

COMPRIMIDO ORAL.

ANTIPARASITARIO INTERNO DE AMPLIO ESPECTRO

Technical Specification

SPECIES

Cats.

Broad spectrum internal antiparasitic, effective in the treatment and control of gastrointestinal nematodes and tapeworms in cats, such as *Toxacara catti*, *Uncinaria stenocephala, Ancylostoma tubaeforme, Taenia taeniformis* and *Dipylidium caninum*.

DOSAGE FORM

Oral tablet.

THERAPEUTIC ACTION

Broad spectrum internal antiparasitic

COMPOSITION

Each tablet contains:
Pyrantel Pamoate ..... 230 mg
(Equivalent to 79.79 mg of Pyrantel)
Praziquantel ..... 20 mg
Excipients q.s. ..... 1 tablet

INDICATIONS

- Do not use in kittens less than 1 month old or weighing less than 700 g
- Do not administer to pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer directly into the oral cavity or incorporated in a small amount of food. It does not require fasting.
Dosage of active ingredients:
Praziquantel 5 mg / Kg of weight in a single dose.
Pyrantel: 20 mg / kg of body weight in a single dose.

Product dosage:
1 tablet every 4 Kg of weight in a single dose.

Cat weight	Dose
1 Kg	1/4 tablet
2 Kg	1/2 tablet
3 Kg	3/4 tablet
4 Kg	1 tablet

DRUG INTERACTIONS

Do not administer simultaneously with Piperazine or Levamisole.

CONTRAINDICATIONS

- Do not use in kittens less than 1 month old or weighing less than 700 g
- Do not administer to pregnant or lactating females.

PRECAUTIONS

- Do not administer simultaneously with Piperazine or Levamisole.
- Use with caution in patients with impaired liver function.

SPECIAL PRECAUTIONS FOR USE

- Do not administer simultaneously with Piperazine or Levamisole.
- Use with caution in patients with impaired liver function.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of water. Dispose of the waste of this product with care together with household waste.

CONSERVATION

Store in a cool and dry place, at room temperature between 15 ° and 30 ° C, protected from light.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

Container case with 15 unit cases.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N°: 1386
Perú: Reg. SENASA F.08.21.I.1138

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Peru by Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

NAXPET® INJECTABLE SOLUTION.

SOLUCIÓN INYECTABLE.

ANALGÉSICO, ANTIPIRÉTICO Y ANTIINFLAMATORIO NO ESTEROIDAL.

Technical Specification

SPECIES

Dogs and cats.

Indicated for the treatment of feverish symptoms, inflammations and painful conditions of the bones, joints and musculoskeletal system in dogs and cats. Its use is especially recommended for the treatment of post-surgical pain, and in cases of osteoarthritis, inflammation of the musculoskeletal system and traumatic injuries in general.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Analgesic, Antipyretic and NON-STEROIDAL Anti-inflammatory.

COMPOSITION

Each mL of solution for injection contains:
Ketoprofen ..... 10 mg
Excipients q.s.p ..... 1 mL

INDICATIONS

- Do not use in animals with known hypersensitivity to Ketoprofen.
- Do not use in animals with pathologies that present with gastrointestinal ulceration or bleeding.
- Do not use concomitantly with anticoagulants or corticosteroids.
- Do not use in animals with severe liver and kidney failure.
- Do not use concomitantly with other NSAIDs, or with diuretics.
- Do not use in pregnant females or in lactation period.

ROUTE OD ADMINISTRATION AND DOSAGE

Route of administration:

- **Dogs:** Subcutaneous, intramuscular or intravenous route.
- **Cats:** Subcutaneous route.

Dose:
Dogs and cats:
0.2 mL per Kg of weight (2 mg / Kg), once a day, for up to 3 consecutive days.

CONTRAINDICATIONS

- Do not use in animals with known hypersensitivity to Ketoprofen.
- Do not use in animals with pathologies that present with gastrointestinal ulceration or bleeding.
- Do not use concomitantly with anticoagulants or corticosteroids.
- Do not use in animals with severe liver and kidney failure.
- Do not use concomitantly with other NSAIDs, or with diuretics.
- Do not use in pregnant females or in lactation period.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

In some patients, edema, nausea, gastrointestinal irritation, headaches, nervousness, constipation, hives, skin rash or fever may be observed.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:

- Discard any unused product remains in its original container.
- Dispose of the waste of this product with care together with household waste.

CONSERVATION

Store in a cool and dry place, at room temperature between 15 and 30 ° C.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

20 mL vial

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 0638
Uruguay: Reg. N° MGAP A-4441
Rep. Dominicana: Reg. Nº 8862
Perú: Registro SENASA F.99.42.I.0075

NAXPET® LARGE BREED - ORAL TABLET

COMPRIMIDO ORAL.

ANALGÉSICO, ANTIPIRÉTICO Y ANTIINFLAMATORIO NO ESTEROIDAL.

Technical Specification

SPECIES

Dogs over 30 Kg.

Analgesic, antipyretic and non-steroidal anti-inflammatory.

DOSAGE FORM

Oral tablet.

THERAPEUTIC ACTION

Analgesic, antipyretic and non-steroidal anti-inflammatory.

COMPOSITION

Each tablet contains:
Ketoprofen ..... 30 mg
Excipients q.s.p ..... 1 comp.

INDICATIONS

- Do not use in case of hypersensitivity to Ketoprofen
- Do not administer to pregnant or lactating females.
- Do not administer in dogs with gastric ulcers
- Do not use simultaneously with anticoagulants or diuretics.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration way: Oral.

- **Dosage**: 1 tablet. / 30 Kg of b.w. (1 mg of Ketoprofen / Kg b.w.) every 24 hours for 5 days. In case of postoperative pain and chronic pain, administer an initial dose of 2 tablets / 30 kg bw (2 mg of Ketoprofen / Kg of b.w.) and then 1 tablet / 30 Kg of b.w. (1 mg of Ketoprofen / Kg of b.w) every 24 hours.

CONTRAINDICATIONS

- Do not use in case of hypersensitivity to Ketoprofen
- Do not administer to pregnant or lactating females.
- Do not administer in dogs with gastric ulcers
- Do not use simultaneously with anticoagulants or diuretics.

PRECAUTIONS

Simultaneous use with other NSAIDs or with corticosteroids can increase side effects, such as vomiting, diarrhea and transient anorexia in dogs.

SPECIAL PRECAUTIONS FOR USE

Simultaneous use with other NSAIDs or with corticosteroids can increase side effects, such as vomiting, diarrhea and transient anorexia in dogs.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Do not handle by people who are hypersensitive to Ketoprofen.
In case of contact with skin, eyes or mucous membranes, wash immediately with plenty of water. In case of accidental ingestion, go immediately to a medical center and show the product box.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

In certain cases it can cause vomiting, diarrhea and transient anorexia. Edema, nausea, gastrointestinal irritation and ulceration, nervousness, constipation, hives, skin rush, or fever can also be observed. These symptoms disappear quickly when treatment is stopped.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 ° C, protected from light.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

Box with 10 tablets.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 1385
Rep. Dominicana: Reg. Nº 8864
El Salvador: Reg. Nº VE2014034895
Perú: Reg. SENASA F.99.21.I.0107
Costa Rica: Reg. N° MV-6986

COUNTRIES WHERE IT IS MARKETED

Distribution in El Salvador:
Rafael Alfredo Alfaro Castillo.
8th C. Pte. Pje. Moreno N ° 112, Col. Flor Blanca.
San Salvador, El Salvador.

Imported and Distributed in Peru by:
Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

NAXPET® ORAL SUSPENSION

SUSPENSIÓN ORAL.

ANALGÉSICO, ANTIPIRÉTICO Y ANTIINFLAMATORIO NO ESTEROIDAL.

Technical Specification

SPECIES

Dogs and cats.

Indicated for the treatment of feverish pictures, inflammations and painful conditions of the bones, joints and skeletal muscle system in dogs and cats. Its use is especially recommended for the treatment of post-surgical pain, and in cases of osteoarthritis, inflammation of the musculoskeletal system and traumatic injuries in general.

DOSAGE FORM

Oral suspension

THERAPEUTIC ACTION

Analgesic, Antipyretic and Non-steroidal anti-inflammatory

COMPOSITION

Each 100 mL of suspension contains:
Ketoprofen ..... 400 mg
Excipients q.s.p ..... 100 mL

INDICATIONS

- Do not use in animals with known hypersensitivity to Ketoprofen.
- Do not use in animals with pathologies that present with ulceration or gastrointestinal bleeding.
- Do not use in animals with severe kidney and liver failure.
- Do not use concomitantly with anticoagulants or corticosteroids.
- Do not use concomitantly with other NSAIDs.
- Do not use in pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration route:
Oral administration.

Dose:
1 mg / Kg once a day for 5 days, (equivalent to 1 mL for every 4 kilos of weight or 8 drops per Kg of weight, once a day for 5 days).
In the case of post-operative pain and chronic pain, administer an initial dose of 2 mg / Kg and then 1 mg / Kg once a day for 5 days (equivalent to an initial dose of 2 mL every 4 Kg of weight or 16 drops per Kg of weight and then 1 mL every 4 Kg of weight or 8 drops per Kg of weight, once a day for 5 days).

CONTRAINDICATIONS

- Do not use in animals with known hypersensitivity to Ketoprofen.
- Do not use in animals with pathologies that present with ulceration or gastrointestinal bleeding.
- Do not use in animals with severe kidney and liver failure.
- Do not use concomitantly with anticoagulants or corticosteroids.
- Do not use concomitantly with other NSAIDs.
- Do not use in pregnant or lactating females.

PRECAUTIONS

Warning and special precautions for use:

- To avoid the risk of increased adverse effects in dogs and cats, it is recommended not to administer the formulation in conjunction with other non-steroidal anti-inflammatory drugs.
- Ketoprofen should be used with caution in animals that present gastrointestinal pathologies or chronic inflammatory bowel disease, and should not be used if there is gastric ulceration or gastrointestinal bleeding.
- Avoid using the formulation in dehydrated, hypovolemic, hypotensive or hypoproteinemic animals, since there is a greater potential risk of kidney toxicity in them.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR USE

Warning and special precautions for use:

- To avoid the risk of increased adverse effects in dogs and cats, it is recommended not to administer the formulation in conjunction with other non-steroidal anti-inflammatory drugs.
- Ketoprofen should be used with caution in animals that present gastrointestinal pathologies or chronic inflammatory bowel disease, and should not be used if there is gastric ulceration or gastrointestinal bleeding.
- Avoid using the formulation in dehydrated, hypovolemic, hypotensive or hypoproteinemic animals, since there is a greater potential risk of kidney toxicity in them.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- After handling and applying the product, a hand wash is recommended.
- Avoid contact of the product with the skin and with the eyes.
- In case of accidental ingestion of the product, it is recommended to go to a medical center and show the internal brochure of the product.

WARNINGS

ADVERSE EFFECTS

Ketoprofen can cause gastrointestinal disturbances at therapeutic doses in some animals. It can cause vomit, diarrhea and transient anorexia in dogs and cats.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
Empty containers can be discarded as household waste, without any special precautions. Do not dispose of containers with product residues on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

- Store in a dry place protected from light, at room temperature between 15 and 30 ° C.
- Once opened, use the product within 3 months. Discard the unused product after that period of time.

CONDITION OF SALE

Sale with a Veterinary Medical prescription only .

PRESENTATION

20 mL bottle

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 1995
Costa Rica: Reg. N° MAG CL 4-14-4-6177
Uruguay: Reg. MGAP Nº 2018A00608
Perú: Registro SENASA F.C6.02.I.0001

COUNTRIES WHERE IT IS MARKETED

Importer in Uruguay:
VIVAFIL S.A.
Rio Negro 1107 Montevideo - Uruguay, TEL 29001112
grupotecnovet@gmail.com
Technical Director: DMTV Diego Cuadrado.

Imported and Distributed in Peru by:
Representaciones Durand SAC.
Av. Manuel Olguín N ° 501 Office N ° 604 Santiago de Surco Lima.

NAXPET® ORAL TABLET

COMPRIMIDO ORAL.

ANALGÉSICO, ANTIPIRÉTICO Y ANTIINFLAMATORIO NO ESTEROIDAL.

Technical Specification

SPECIES

Dogs and cats

Analgesic, Antipyretic and Non-steroidal Anti-inflammatory.

DOSAGE FORM

Oral tablet

THERAPEUTIC ACTION

Analgesic, Antipyretic and Non-steroidal Anti-inflammatory.

COMPOSITION

Each tablet contains:
Ketoprofen ..... 10 mg
Excipients q.s.p ..... 1 comp.

INDICATIONS

Do not use in pregnant or lactating females; in animals with gastrointestinal ulcers, severe liver or kidney damage; in animals hypersensitive to Ketoprofen or simultaneously with anticoagulants or diuretics.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration way: Oral.

Dose: 1 comp./10 kg (1 mg / Kg) every 24 hours for 5 days. In the case of postoperative pain and chronic pain, administer an initial dose of 2 comp./10 Kg (2 mg / Kg) and then 1 comp./10 Kg (1 mg / Kg) every 24 hours.

DRUG INTERACTIONS

- Due to its high affinity for plasma proteins, Ketoprofen can displace or be displaced by other drugs with the same characteristics.
- Ketoprofen may increase the likelihood of gastroenteric bleeding or ulceration if used concomitantly with other medicinal products that alter hemostasis and / or cause intestinal erosion.

CONTRAINDICATIONS

Do not use in pregnant or lactating females; in animals with gastrointestinal ulcers, severe liver or kidney damage; in animals hypersensitive to Ketoprofen or simultaneously with anticoagulants or diuretics.

SPECIAL PRECAUTIONS FOR THE OPERATOR

After handling and applying the product, a hand wash is recommended. Avoid contact of the product with the skin and with the eyes. In the event of accidental ingestion of the product, it is recommended to go to a healthcare center.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
Discard the remains of unused product in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of natural water. Dispose of this product with caution with household waste.

CONSERVATION

Store in a cool, dry place, at room temperature between 15 and 30 ° C.

CONDITION OF SALE

Sale with a Veterinary Medical prescription.

PRESENTATION

Case with 10 tablets.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 0583
Rep. Dominicana: Reg. Nº 9304
Uruguay: Reg. MGAP N° A-4458
Perú: Registro SENASA F.99.21.I.0074

COUNTRIES WHERE IT IS MARKETED

Uruguay:
Importador: VIVAFIL S.A.
Rio Negro 1107 Montevideo - Uruguay, TEL 29001112
grupotecnovet@gmail.com
Director Técnico: DMTV Diego Cuadrado.

Perú:
Importado y Distribuido por Representaciones Durand SAC.
Av. Manuel Olguín N° 501 Oficina N° 604 Santiago de Surco Lima.

Bolivia:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

OFTAVET® OPHTHALMIC SOLUTION

SOLUCIÓN OFTÁLMICA.

ANTIBIÓTICO TÓPICO OFTÁLMICO.

Technical Specification

SPECIES

Dogs, cats and horses.

Oftavet® Ophthalmic Solution is an antibiotic for topical use indicated to treat superficial and deep ocular infections of the eye and its annexes, including corneal ulcers infected with bacteria sensitive to the drug.

DOSAGE FORM

Ophthalmic solution

THERAPEUTIC ACTION

Ophthalmic topical antibiotic.

COMPOSITION

Each 1 mL of product contains:
Ciprofloxacin Hydrochloride Monohydrate ..... 3.49 mg
(Equivalent to 0.3% of Ciprofloxacin base)
Anhydrous Disodium EDTA ..... 2.33 mg
Excipients q.s.p ..... 1 mL

PROPERTIES

Ciprofloxacin is a Fluoroquinolone with bactericidal action. Its action is based on the inhibition of bacterial DNA gyrase or topoisomerase IV (a type of topoisomerase II), thereby preventing DNA supercoiling and replication. Respiration and cell division are terminated, and other processes are disrupted, including membrane integrity. Ciprofloxacin has a wide spectrum of activity, being highly effective against the most commonly found ocular pathogens, both Gram positive and Gram negative, such as: Staphylococcus spp., Streptococcus spp., Pseudomonas aeruginosa, Pasteurella multocida, E. coli, etc.

EDTA inhibits metalloproteinases by chelation of Zinc and Calcium that the enzyme requires as a cofactor and a stabilizing ion, respectively. By chelating the calcium ion, EDTA interferes with the stability of metalloproteinases and thus decreases the stimulation for the migration of polymorphonuclear leukocytes to the corneal ulcer site. EDTA also interferes with the adhesion of metalloproteinases to the polymorphonuclear cell membrane, leading the cell membrane to an inactivated state. EDTA exhibits a high rate of anti-metalloproteinase activity in vitro (99.4%).

INDICATIONS

Oftavet® Ophthalmic Solution is contraindicated in patients with hypersensitivity to active ingredients or other quinolones.
Do not use in pregnant or lactating females.

Use during pregnancy and lactation:
Because there is no history about the use of Ciprofloxacin via the eye in pregnant females, it is recommended not to use in this state, nor during lactation.

ROUTE OD ADMINISTRATION AND DOSAGE

Topical administration by instillation into the lower ocular sac.
Administer aseptically. Wash your hands before handling and avoid the bottle coming into contact with the patient's eyes.

- **Horses:** 2 drops every 6 hours for 7 days.
- **Dogs and cats:** 1 drop every 6 hours for 7 days.

DRUG INTERACTIONS

Synergism can occur with aminoglycosides, 3rd generation cephalosporins, and broad spectrum penicillins.

CONTRAINDICATIONS

Oftavet® Ophthalmic Solution is contraindicated in patients with hypersensitivity to active ingredients or other quinolones.
Do not use in pregnant or lactating females.

Use during pregnancy and lactation:
Because there is no history about the use of Ciprofloxacin via the eye in pregnant females, it is recommended not to use in this state, nor during lactation.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- To reduce the risk of contact, people who are hypersensitive to quinolones should wear rubber gloves to avoid contact.
- In case of skin contact, immediately wash the affected area with plenty of water. If irritation persists, consult a doctor.
- In case of eye contact, wash immediately with plenty of water, if vision is blurred, immediately take to a medical center.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

A hypersensitivity could cause erythema, itching, swelling, tearing. In the event of any hypersensitivity reaction, discontinue use and consult a Veterinarian.

Security:
The use of ophthalmic Ciprofloxacin administered in concentrations of 0.3% and 0.75% for one month in Beagle puppies does not produce any local or systemic adverse effects.

GUARD PERIOD

Do not administer to horses intended for human consumption.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Do not dump the empty container or with product remains in rivers, lakes or streams of natural water. Dispose of this product waste carefully with household waste. Contact the manufacturing company or companies specialized in the elimination of waste, to receive recommendations on the disposal of expired or unused products.

CONSERVATION

Keep at a temperature between 2 and 30 ° C and protected from light.
Uruguay: The validity period once the container is opened is 4 weeks.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

Dropper bottle with 5 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG Nº 2105
Bolivia: Reg. SENASAG PUV-F-N° 005509/13
Uruguay: Reg. MGAP Nº 2018A00614

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

Importer in Uruguay:

VIVAFIL S.A.
RIO NEGRO 1107 Montevideo - Uruguay, TEL 29001112
grupotecnovet@gmail.com
Technical Director: DMTV Diego Cuadrado.

OSTEODRAG HA® TABLETS

COMPRIMIDO ORAL.

SUPLEMENTO NUTRICIONAL PARA PERROS.

Technical Specification

SPECIES

Dogs of all ages.

Advanced formula for dogs with joint problems.

DOSAGE FORM

Tablets.

THERAPEUTIC ACTION

Nutritional supplement for dogs.

COMPOSITION

Each tablet contains:

Glucosamina Sulfato.....500 mg

Condroitín Sulfato.....400 mg

Metilsulfonilmetano (MSM).....250 mg

Colágeno.....130 mg

Vitamina C.....33 mg

Ácido Hialurónico.....15 mg

Manganeso Sulfato.....5 mg

Excipientes c.s.p.....1 comprimido

INGREDIENTS

Glucosamine Sulfate (Crab Shell), Chondroitin Sulfate (Shark Cartilage), Methylsulfonylmethane, Palatable Agent (Hydrolyzed Poultry Offal), Hydrolyzed Collagen (Bovine Leather), Magnesium Stearate, Vitamin C, Hyaluronic Acid, Manganese Sulfate and authorized preservatives.

PROXIMATE ANALYSIS

Guaranteed analysis:

Crude protein (min.) 25.3%

Crude Fat (min) 0.4%

Crude fiber (max) 0.3%

Humidity (max) 16.4%

PROPERTIES

Osteodrag HA® is a nutritional supplement for dogs with nutritional requirements special, which provides advanced support for cushioning and joint mobility. osteodrag HA® provides the adequate nutrients and in optimal quantities for the correct maintenance and repair of articular cartilage and connective tissue. Its use is helpful in different processes osteoarticular that require protection and repair.
Glucosamine Sulfate together with Chondroitin Sulfate, stimulate the synthesis of components structures, promote repair mechanisms and maintain joint viscosity, generating greater strength, flexibility and joint protection. MSM, an organic compound that provides Sulfur, allows an adequate enzymatic, immunological and tissue formation function connective. Collagen is a structural component of the extracellular matrix and articular cartilage. and the Hyaluronic Acid present in the synovial fluid, has lubricating and viscoelastic properties. Vitamin C has antioxidant properties and stimulates the production of Collagen and the Manganese is an essential cofactor in the synthesis of glycosaminoglycans.

USE INSTRUCTIONS

Administer orally according to the following table:

Weight (Kg)	Initial dose (First 6 weeks)	Maintenance dose
5 a 10	1 tablet / day	1/2 tablet / day
11 a 22	2 tablet / day	1 tablet / day
23 a 45	3 tablet / day	1 1/2 tablet /day
> 45	4 tablet / day	2 tablet / day

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

SUPPLEMENT
VETERINARY USE
EXCLUSIVE USE IN ANIMAL FEEDING
IT DOES NOT CORRESPOND TO A COMPLETE FOOD
FORBIDDEN ITS USE IN THE FEEDING OF RUMINANTS

CONSERVATION

Store in a cool, dry place protected from light, at no more than 30ºC.

CONDITION OF SALE

Free sale.

PRESENTATION

30 tablets.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.
Lautaro N° 300. Quilicura. Santiago. Chile.
www.dragpharma.cl

RECORDS

Reg. LENAA N°: RM 03-008N

OTIBACT® OTIC GEL.

GEL ÓTICO.

ANTIBACTERIANO - ANTIMICÓTICO – ANTIPRURIGINOSO - ANTIINFLAMATORIO

Technical Specification

SPECIES

Dogs.

OTIBACT® Otic Gel is indicated for the treatment of acute and chronic external otitis caused by mixed infection of Staphylococcus intermedius or Pseudomonas aeruginosa and Malassezia pachydermatis, sensitive to the active principles of the association. Its anti-inflammatory effect reduces the levels of erythema, itching and secretion in the external auditory canals of dogs.

DOSAGE FORM

Otic Gel.

THERAPEUTIC ACTION

Antibacterial - Antifungal - Antipruritic - Anti-inflammatory

COMPOSITION

Each 5 g syringe contains:
Ciprofloxacin Hydrochloride Monohydrate
(Equivalent to 50 mg of Ciprofloxacin base) ..... 58.2 mg
Clotrimazole ..... 50.0 mg
Triamcinolone Acetonide ..... 2.5 mg
Excipients csp ..... 5.0 g

INDICATIONS

- Contraindicated in patients with hypersensitivity to any of its components.
- Do not treat animals with perforation of the tympanic membrane.
- Do not administer in females during the gestation or lactation period.
- Do not administer in breeding animals.
- Do not use with other drugs that can induce ototoxicity.
- Do not use in dogs under 6 months of age and / or with less than 1 kg of body weight.

MODE OF APPLICATION

Clean and dry the external auditory canal, removing wax and exudate. Trim excess hair from the area to be treated. Apply a dose of 0.5 g (one mark of the graduated syringe), once a day, directly into the ear canal, for 7 to 10 consecutive days. One syringe contains 10 doses for the complete treatment of one ear. The plunger of the syringe is graduated with marks from 1 to 10, each of which allows a dose of 0.5 g to be administered. Turn the wheel that comes at the base of the plunger, retracting it towards the free end of the plunger and fix it at the mark corresponding to the day of treatment. Once the wheel is set on the mark corresponding to the day of administration, push the plunger until the wheel stops against the base of the syringe to release the contents. After application, massage the ear canal for a better distribution of the product.

ROUTE OD ADMINISTRATION AND DOSAGE

Topical administration in the external auditory canal.

DRUG INTERACTIONS

Do not apply in conjunction with another substance or veterinary drug.

CONTRAINDICATIONS

- Contraindicated in patients with hypersensitivity to any of its components.
- Do not treat animals with perforation of the tympanic membrane.
- Do not administer in females during the gestation or lactation period.
- Do not administer in breeding animals.
- Do not use with other drugs that can induce ototoxicity.
- Do not use in dogs under 6 months of age and / or with less than 1 kg of body weight.

PRECAUTIONS

- Before starting treatment, the identification of the etiological agent (s) should be carried out, either by smear or culture. In addition, the antibiotic susceptibility of pathogenic bacteria should be evaluated before using this preparation, since fluoroquinolones, including Ciprofloxacin, should not be used as the first line of treatment, unless there is justification for their use.
- There is evidence that corticosteroids can be absorbed after topical application, causing systemic effects. Consequently, any animal using this product should be closely watched for signs such as polydipsia, polyuria, or increased weight gain.
- Before applying the product, the external auditory canal should be examined to ensure the integrity of the tympanic membrane (in order to prevent damage to the vestibular and cochlear apparatus).

SPECIAL PRECAUTIONS FOR USE

- Before starting treatment, the identification of the etiological agent (s) should be carried out, either by smear or culture. In addition, the antibiotic susceptibility of pathogenic bacteria should be evaluated before using this preparation, since fluoroquinolones, including Ciprofloxacin, should not be used as the first line of treatment, unless there is justification for their use.
- There is evidence that corticosteroids can be absorbed after topical application, causing systemic effects. Consequently, any animal using this product should be closely watched for signs such as polydipsia, polyuria, or increased weight gain.
- Before applying the product, the external auditory canal should be examined to ensure the integrity of the tympanic membrane (in order to prevent damage to the vestibular and cochlear apparatus).

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Wash your hands after administering the product.
- If the product comes into contact with the eyes, wash with plenty of water.
- In the case of accidental ingestion, do not induce vomiting. Get medical help.
- Do not handle the drug if you have a known hypersensitivity to any of its components.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

- Occasionally Clotrimazole may cause erythema, hives, blisters, edema, itching, and generally skin irritation.
- In dogs, Cushing's syndrome has been reported in association with prolonged and repeated therapy with corticosteroids.
- Using this medicine for more than 10 days may delay wound healing.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Dispose of any unused product remains in its original, tightly closed container. Dispose of this product waste carefully with household waste. Contact the manufacturing company or companies specialized in waste disposal, to receive recommendations on the disposal of expired or unused products.

CONSERVATION

Store at a temperature between 2 and 30 ° C, protected from light.
Once opened, the syringe should be used within 10 days and stored between 15 and 30ºC, protected from light.

CONDITION OF SALE

Chile: Sale under Veterinary Medical prescription restricted.
Peru: Free Sale.

PRESENTATION

1 syringe with 5 g of product

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG Nº 2322
Perú: Reg. SENASA F.F9.66.I.0008

OVOLUTE® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

PROSTAGLANDINA SINTÉTICA.

Technical Specification

SPECIES

Bovines, sheep, horses and pigs.

1. **Induction or synchronization of estrus.**
   If a functional corpus luteum is diagnosed, injection of Ovolute® causes luteolysis and oestrus with ovulation between the 2nd and 4th day after administration.
   The corpus luteum or cysts disappear within a few days. Ovolute® can also be used to shorten the cycle or timing of estrus.
2. **Physiological or pathological expulsion of uterine contents.**
   Due to the stimulating effect of Ovolute® on the muscles of the uterus, the contents of the uterus are expelled. This effect can be used for the induction of parturition (not before 260 days in cattle and 112 days in pigs), abortion (before day 150 in cattle and 35 in horses) and in endometritis. Abortion or expulsion usually occurs on the 1st or 2nd day after the injection.

DOSAGE FORM

Injectable solution

THERAPEUTIC ACTION

Synthetic prostaglandin

COMPOSITION

Each 100 mL of solution for injection contains:
Cloprostenol Sodium ..... 7.89 mg
(Equivalent to 7.5 mg D (+) - Cloprostenol base)
Excipients q.s.p ..... 100 mL

PROPERTIES

D-Cloprostenol is a synthetic analog of PGF2a and has a biological activity similar to this natural endogenous prostaglandin. D-Cloprostenol is capable of producing the luteolytic and myometrial-stimulating effects of PGF2 Alpha in small doses, which is the basis for its therapeutic indications.

INDICATIONS

- Do not use in pregnant females, except for the indicated indications (abortion or childbirth).
- Do not administer to animals hypersensitive to Cloprostenol.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration by intramuscular route.

- **Cattle:** 2 mL (equivalent to 150 µg of D-Cloprostenol), in all its indications.
  Induction of oestrus and ovulation: From the 2nd day after administration, animals should receive adequate supervision of oestrus.
  Oestrus timing: repeat dose 11 days later. Estrus occurs 2 to 4 days later.
- **Horses:** 0.3 to 1 mL (equivalent to 25 to 75 µg of D-Cloprostenol), in all its indications.
  Induction of estrus and shortening of the cycle.
  In labor induction, administer after day 320 of gestation. Delivery usually takes a few hours.
- **Pigs:** 1 mL (equivalent to 75 µg of D-Cloprostenol), in all its indications.
  Induction of labor: Administer after day 112 of gestation. In 70% of cases, delivery should occur at 19 and 30 hours after treatment.
- **Sheep:** 1 mL (equivalent to 75 µg of D-Cloprostenol).
  To synchronize estrous with progestogens, apply a single dose after removal of the intravaginal device.
  To synchronize estrus only with prostaglandins, apply two doses with an interval of 7 to 9 days. Estrus occurs 2 to 4 days after the second dose.

DRUG INTERACTIONS

Do not use in animals under treatment with non-steroidal anti-inflammatory drugs, since the synthesis of endogenous prostaglandins.
The activity of other oxytocic agents may be increased after cloprostenol administration.

CONTRAINDICATIONS

- Do not use in pregnant females, except for the indicated indications (abortion or childbirth).
- Do not administer to animals hypersensitive to Cloprostenol.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Avoid contact with pregnant women and people suffering from diseases of the tract respiratory (such as asthma).
- Do not handle by people who are hypersensitive to Cloprostenol.
- Avoid contact with skin; in case of contamination wash with plenty of water.
- In case of accidental injection, go immediately to a medical center and show the label of the product.
- Do not smoke, eat or drink during the administration of the product.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Meat: 1 day.

Milk: 0 days.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:

- Empty containers can be disposed of as household waste, without any special precautions.
- Do not dispose of containers with product remains on the ground or water courses.
- For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 ° C, protected from light. Once the container is opened, use within 12 weeks.
Discard the unused product after that period of time.

CONDITION OF SALE

Sale under retained Veterinary Medical prescription.

PRESENTATION

20 mL and 50 mL vial

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Reg. SAG Nº 1600

PACIFOR® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

SEDANTE - PRE ANESTÉSICO

Technical Specification

SPECIES

Dogs

Pacifor® Solution for Injection, is indicated as a sedative and pre anesthetic agent.

DOSAGE FORM

Injectable solution

THERAPEUTIC ACTION

Sedative - Pre-anesthetic

COMPOSITION

Each 1 mL of solution for injection contains:
Accepromazine Maleate ..... 10 mg
(Equivalent to 7.4 mg of acepromazine base)
Excipients q.s.p ..... 1 mL

INDICATIONS

- Do not administer intra-arterially.
- Do not administer to animals with hypovolemia or shock.
- Do not administer to animals with coagulopathies or thrombocytopenia.
- Do not administer to animals with tetanus or strychnine poisoning.
- Do not administer within one month of deworming with Organophosphate agents.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration routes:
Intramuscular, intravenous or subcutaneous.

Dose: 0.14 - 0.27 mL per 10 kilos of body weight in a single dose, equivalent to 0.1 - 0.2 mg of acepromazine base per kilo of body weight, in a single dose. Do not exceed 3 mg total dose.

After parenteral administration, animals should be kept in a quiet place until the onset of sedative effects that occur around 15 minutes. The effects can last up to 3 hours. Endovenous administration should be slow. Administer aseptically.

DRUG INTERACTIONS

- Accepromazine may exacerbate the action of central nervous system depressants if used together.
- Concomitant administration with Epinephrine can lead to beta activity causing vasodilation and increased heart rate.
- Opioids can improve the hypotensive effects of Accepromazine.
- Accepromazine should not be administered within one month of deworming with organophosphate agents as its effects may be potentiated.
- Phenytoin metabolism may decrease if given simultaneously with phenothiazines.
- Procaine activity can be enhanced by phenothiazines.
- Co-administration of Acepromazine and Propanolol can lead to serum elevation of both drugs.
- Co-administration of Quinidine with phenothiazines can cause additive cardiac depression.
- Vasocontrictors like Phenylephrine antagonize the hypotensive effects of phenothiazines.

CONTRAINDICATIONS

- Do not administer intra-arterially.
- Do not administer to animals with hypovolemia or shock.
- Do not administer to animals with coagulopathies or thrombocytopenia.
- Do not administer to animals with tetanus or strychnine poisoning.
- Do not administer within one month of deworming with Organophosphate agents.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- In the case of ocular exposure, the eyes should be irrigated with plenty of water. If irritation exists and persists, the patient should be evaluated by a doctor.
- In the case of dermal exposure, contaminated clothing should be removed or the exposed area washed thoroughly with water. If irritation exists and it persists, the patient should be evaluated by a doctor.
- In case of accidental ingestion, the mouth should be washed. If there was large intake, immediate medical attention should be requested.
- In the case of accidental injection, immediate medical attention should be requested.

WARNINGS

Advertencias y precauciones especiales de uso:

- La administración endovenosa debe realizarse en forma lenta.
- Luego de la administración de Acepromazina, se requieren menores dosis de anestésicos generales.
- Usar con precaución en procedimientos anestésicos regionales con anestésicos locales.
- Utilizar con precaución y con la menor dosis recomendada en animales con disfunción hepática, enfermedad cardíaca y debilidad general.
- Utilizar con precaución en animales muy jóvenes o debilitados.
- Usar con precaución en pacientes geriátricos.
- Acepromazina no posee efectos analgésicos, se deben tomar medidas para controlar el dolor.
- Usar con precaución en perros braquicéfalos (Bóxer, Pequinés, etc.), en perros con mutaciones MDR1 (Collies, Pastores australianos, entre otros) y en razas gigantes, Galgos y Terrier.
- Usar con precaución como agente de contención en perros agresivos.
- Se puede utilizar en conjunto con Atropina para bloquear sus efectos bradicárdicos.
- Se recomienda utilizar lubricantes oculares o lágrimas artificiales como protectores corneales.
- Una dosis mayor no acelera el inicio de la sedación, solo alarga el efecto sedante.
- Una vez que se ha administrado, es importante dejar al animal en un lugar tranquilo por alrededor de 15 minutos o hasta que se logren los efectos sedantes.
- Luego de la administración, los pacientes no deben estar expuestos a fluctuaciones de temperaturas extremas por al menos 8 horas.
- La administración en conjunto con otros depresores respiratorios o del Sistema Nervioso Central, puede exacerbar la depresión respiratoria.
- Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

Unwanted effects and adverse reactions:

- Accepromazine generates vasodilation-mediated hypotension, which can last for a few hours. It is usually mild in healthy animals, however, it can be accentuated in anesthetized, weakened or hypovolemic patients.
- Unusual cardiovascular collapse of varying severity (secondary to bradycardia and hypotension) has been described, especially in brachycephalic breeds.
- Accepromazine causes significant intraoperative hypothermia that lasts for a few hours.
- Accepromazine causes prolapse of the nictitating membrane, which lasts for a few hours.
- Accepromazine may transiently decrease hematocrit, reestablishing itself after several hours, which may be of importance in anemic animals.
- Accepromazine transiently decreases platelet aggregation without affecting clotting times.
- Occasionally Accepromazine can generate contradictory clinical signs of aggressiveness and generalized stimulation of the Central Nervous System. These reactions are infrequent and transient in nature and generally do not last for more than 48 hours.
- Intramuscular injections of Acepromazine can cause transient pain at the injection site.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
Discard the unused product in its original container.
Dispose of empty containers of this product along with household waste.
Contact the manufacturing company or companies specialized in waste disposal, to receive recommendations on the disposal of expired or unused products.

CONSERVATION

Store at a temperature between 15 and 30 ° C, away from light. Once the container is opened, use within 3 months and it must be stored at a temperature between 15 and 30ºC, sheltered from light.

PRESENTATION

Ampoule bottle with 10 mL and 50 mL

RECORDS

Chile: SAG Reg. No. 2390
Bolivia: Reg. SENASAG PUV-F-N° 005506/13
Imported and distributed by:
ZOO PHARMA VETERINARY INPUTS S.R.L.
Diaz Romero N° 1339 Miraflores Zone, La Paz-Bolivia
Telephone: 591 2223357
Peru: Reg. SENASA F.62.01.I.0028
Imported and distributed by Representaciones Durand
SAC. Av. Manuel Olguín N° 501 Office N° 604 Santiago de Surco Lima.
Costa Rica: Reg. No. MV-7152

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

PACIFOR® ORAL SOLUTION

SOLUCIÓN ORAL.

TRANQUILIZANTE.

Technical Specification

SPECIES

Dogs and cats.

Pacifor® Oral solution, is indicated as a sedative (tranquilizer) agent for dogs and cats.

DOSAGE FORM

Oral solution

THERAPEUTIC ACTION

Tranquilizer

COMPOSITION

Each 100 mL of solution contains:
Acepromazine Maleate ..... 1.35 g
(Equivalent to 1 g of acepromazine base)
Excipients q.s.p... ..... 100 mL

INDICATIONS

- Do not administer to pregnant or lactating females.
- Do not use in animals undergoing tests such as myelograms.
- Do not use in animals with a history of seizures.
- Do not administer in animals with hypovolemia or shock, due to its hypotensive effects.
- Do not administer in animals with coagulopathies or thrombocytopenia, due to its effects on the platelet aggregation.
- Do not administer to animals with tetanus or Strychnine poisoning, due to the effects on the extra pyramid system.
- Do not administer within one month of deworming with Organophosphate agents already that its effects can be enhanced.

USE INSTRUCTIONS

The calming effect takes between 30 and 40 minutes. Place the animal in a calm environment. The duration of the effect is 6 to 12 hours.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral Administration.

Product dosage:

- Dogs: 1 to 2 drops per Kg of weight (0.5 to 1 mg / Kg).
- Cats 2 to 3 drops per Kg of weight (1 to 1.5 mg / Kg).

DRUG INTERACTIONS

- Combinations of antidiarrheals (eg Kaolin-Pectin, mixtures with Bismuth Subsalicylate) and antacids can reduce gastrointestinal absorption of orally administered phenothiazines.
- Depressants of the Central Nervous System (CNS) (barbiturates, narcotics, anesthetics, etc.), can exacerbate CNS depression if used together with Acepromazine.
- Epinephrine: Phenothiazines block alpha-adrenergic receptors; concomitant administration with epinephrine can lead to beta activity causing vasodilation and increased heart rate.
- Opioids: May improve the hypotensive effects of Acepromazine. Acepromazine doses are generally reduced when used with an opiate.
- Organophosphate agents: Acepromazine should not be administered within one month of deworming with these agents as their effects may be potentiated.
- Phenytoin: Your metabolism may decrease if given simultaneously with phenothiazines.
- Procaine: Activity can be enhanced by phenothiazines.
- Propranolol: Co-administration of Acepromazine and Propanolol can lead to serum elevation of both drugs.
- Quinidine: With phenothiazines it can cause additive cardiac depression.
- Vasocontrictors like Phenylephrine antagonize the hypotensive effects of phenothiazines.

CONTRAINDICATIONS

- Do not administer to pregnant or lactating females.
- Do not use in animals undergoing tests such as myelograms.
- Do not use in animals with a history of seizures.
- Do not administer in animals with hypovolemia or shock, due to its hypotensive effects.
- Do not administer in animals with coagulopathies or thrombocytopenia, due to its effects on the platelet aggregation.
- Do not administer to animals with tetanus or Strychnine poisoning, due to the effects on the extra pyramid system.
- Do not administer within one month of deworming with Organophosphate agents already that its effects can be enhanced.

PRECAUTIONS

- After the administration of Acepromazine, lower doses of general anesthetics are required.
- Use with caution and at the lowest recommended dose in animals with liver dysfunction, heart disease and general weakness.
- Use with caution in very young or weakened animals, due to its effects on thermoregulation.
- Use with caution in geriatric patients, as very low doses have been associated with prolonged effects of the drug.
- Acepromazine does not have analgesic effects, so adequate measures should be taken to control pain in treated animals.
- Use with caution in brachycephalic dogs (Boxer, Pekingese, etc.) or cats of the Persian breed, as they can be very sensitive to the bradycardic effects of Acepromazine.
- Use with caution in dogs with MDR1 mutations (Collies, Australian Shepherds, among others), as they can develop a deeper sedation that persists longer than usual.
- Giant breeds and Greyhounds can be extremely sensitive to the drug, while terrier breeds are somewhat resistant to its effects.
- Acepromazine should be used with caution as a containment agent in aggressive dogs, as it may make animals more prone to startle and react to noise or other sensory stimuli.
- Atropine can be used in conjunction with Acepromazine to help block its bradycardic effects.
- During sedation with Acepromazine it is recommended to use eye lubricants or artificial tears as corneal protectors, due to the reduction in tear production.
- A higher dose does not accelerate the onset of sedation, it only lengthens the sedative effect.
- Once it has been administered, it is important to leave the animal in a quiet place for about 15 minutes or until sedative effects are achieved.
- After administration, patients should not be exposed to extreme temperature fluctuations for at least 8 hours.
- Administration in conjunction with other respiratory or central nervous system depressants can exacerbate respiratory depression.

SPECIAL PRECAUTIONS FOR USE

- After the administration of Accepromazine, lower doses of general anesthetics are required.
- Use with caution and at the lowest recommended dose in animals with liver dysfunction, heart disease and general weakness.
- Use with caution in very young or weakened animals, due to its effects on thermoregulation.
- Use with caution in geriatric patients, as very low doses have been associated with prolonged effects of the drug.
- Accepromazine does not have analgesic effects, so adequate measures should be taken to control pain in treated animals.
- Use with caution in brachycephalic dogs (Boxer, Pekingese, etc.) or cats of the Persian breed, as they can be very sensitive to the bradycardic effects of Accepromazine.
- Use with caution in dogs with MDR1 mutations (Collies, Australian Shepherds, among others), as they can develop a deeper sedation that persists longer than usual.
- Giant breeds and Greyhounds can be extremely sensitive to the drug, while terrier breeds are somewhat resistant to its effects.
- Accepromazine should be used with caution as a containment agent in aggressive dogs, as it may make animals more prone to startle and react to noise or other sensory stimuli.
- Atropine can be used in conjunction with Accepromazine to help block its bradycardic effects.
- During sedation with Accepromazine it is recommended to use eye lubricants or artificial tears as corneal protectors, due to the reduction in tear production.
- A higher dose does not accelerate the onset of sedation, it only lengthens the sedative effect.
- Once it has been administered, it is important to leave the animal in a quiet place for about 15 minutes or until sedative effects are achieved.
- After administration, patients should not be exposed to extreme temperature fluctuations for at least 8 hours.
- Administration in conjunction with other respiratory or central nervous system depressants can exacerbate respiratory depression.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- In the case of ocular exposure, the eyes should be irrigated with plenty of water. If irritation exists and persists, the patient should be evaluated by a doctor.
- In the case of dermal exposure, contaminated clothing should be removed or the exposed area washed thoroughly with water. If irritation exists and it persists, the patient should be evaluated by a doctor.
- In case of accidental ingestion, the mouth should be washed. If there was large intake, immediate medical attention should be requested.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

- Accepromazine generates hypotension mediated by vasodilation, which can last for a few hours and must be considered during its use. In healthy animals this hypotension is usually mild, however it can be accentuated in anesthetized, weakened or hypovolemic patients.
- Unusual cardiovascular collapse of varying severity (secondary to bradycardia and hypotension) has been described in some dogs, especially in brachycephalic breeds.
- Accepromazine causes significant intraoperative hypothermia because it depresses the hypothalamic thermoregulatory center and causes cutaneous vasodilation. This effect lasts for a few hours.
- Accepromazine causes prolapse of the nictitating membrane, which lasts for a few hours while maintaining its effect.
- Accepromazine may decrease hematocrit for several hours, as a result of splenic congestion after α1-adrenergic receptor blockage, which may be important in anemic animals. This effect is transitory and is restored after several hours.
- Accepromazine transiently decreases platelet aggregation without affecting clotting times.
- Occasionally Accepromazine can generate contradictory clinical signs of aggressiveness and generalized stimulation of the Central Nervous System. These reactions are infrequent and transient in nature and generally do not last for more than 48 hours.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:

- Discard the remains of unused product in its original container.
- Do not throw the empty container or with product remains, in rivers, lakes or torrents of water.
- Do not reuse the container.
- Dispose of this product with caution with household waste.

CONSERVATION

Store between 15 ° and 30° C, protected from light. Once the container is opened, use within 12 weeks. Discard the unused product after that period of time.

CONDITION OF SALE

Sale with Veterinary Medical prescription.

PRESENTATION

Dropper bottle with 10 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N°: 0516
Bolivia: Reg. SENASAG PUV-F-N° 005517/13

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

PACIFOR® ORAL TABLET

COMPRIMIDO ORAL.

TRANQUILIZANTE.

Technical Specification

SPECIES

Dogs and cats.

Pacifor® tablets is indicated as a sedative (tranquilizer) agent for dogs and cats.

DOSAGE FORM

Oral tablet

THERAPEUTIC ACTION

Tranquilizer

COMPOSITION

Each tablet contains:
Accepromazine Maleate ..... 10 mg
(Equivalent to 7.4mg of acepromazine base)
Excipients q.s.p ..... 1 tablet

INDICATIONS

- Do not use in animals undergoing tests such as myelograms.
- Do not use in animals with a history of seizures.
- Do not administer in animals with hypovolemia or shock, due to its hypotensive effects.
- Do not administer to animals with coagulopathies or thrombocytopenia, due to its effects on platelet aggregation.
- Do not administer to animals with tetanus or Strychnine poisoning, due to the effects on the extra pyramidal system.
- Do not administer within one month of deworming with Organophosphate agents since their effects can be potentiated.
- Do not administer to pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration by oral route.

Dose of the active substance:
Dogs: 0.74 mg of acepromazine base per kilogram of weight.
Cats: 1.48 mg of acepromazine base per kilogram of weight.

Product dosage:
Dogs: 1 tablet / 10 Kg of weight.
Cats: 1 tablet / 5 Kg of weight.

Frequency and duration of treatment:
The doses can be repeated according to the criteria of the treating Veterinary Doctor.

DRUG INTERACTIONS

- Combinations of antidiarrheals (eg Kaolin-Pectin, mixtures with Bismuth Subsalicylate) and antacids can reduce gastrointestinal absorption of orally administered phenothiazines.
- Depressants of the Central Nervous System (CNS) (barbiturates, narcotics, anesthetics, etc.), can exacerbate CNS depression if used together with Acepromazine.
- **Epinephrine:** Phenothiazines block alpha-adrenergic receptors; concomitant administration with epinephrine can lead to beta activity causing vasodilation and increased heart rate.
- **Opioids:** May improve the hypotensive effects of Acepromazine. Acepromazine doses are generally reduced when used with an opiate.
- **Organophosphate agents:** Acepromazine should not be administered within one month of deworming with these agents as their effects may be potentiated.
- **Phenytoin:** Your metabolism may decrease if given simultaneously with phenothiazines.
- **Procaine:** Activity can be enhanced by phenothiazines.
- **Propranolol:** Co-administration of Acepromazine and Propanolol can lead to serum elevation of both drugs.
- **Quinidine:** With phenothiazines it can cause additive cardiac depression.
- Vasocontrictors like **Phenylephrine** antagonize the hypotensive effects of phenothiazines.

CONTRAINDICATIONS

- Do not use in animals undergoing tests such as myelograms.
- Do not use in animals with a history of seizures.
- Do not administer in animals with hypovolemia or shock, due to its hypotensive effects.
- Do not administer to animals with coagulopathies or thrombocytopenia, due to its effects on platelet aggregation.
- Do not administer to animals with tetanus or Strychnine poisoning, due to the effects on the extra pyramidal system.
- Do not administer within one month of deworming with Organophosphate agents since their effects can be potentiated.
- Do not administer to pregnant or lactating females.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- In the case of ocular exposure, the eyes should be irrigated with plenty of water. If irritation exists and persists, the patient should be evaluated by a doctor.
- In the case of dermal exposure, the exposed area should be washed with water. If irritation exists and it persists, the patient should be evaluated by a doctor.
- In case of accidental ingestion, the mouth should be washed. If there was large intake, immediate medical attention should be requested.

WARNINGS

Advertencias y precauciones especiales de uso:

- Luego de la administración de Acepromazina, se requieren menores dosis de anestésicos generales.
- Utilizar con precaución y con la menor dosis recomendada en animales con disfunción hepática,enfermedad cardíaca y debilidad general.
- Utilizar con precaución en animales muy jóvenes o debilitados, debido a sus efectos sobre la termorregulación.
- Usar con precaución en pacientes geriátricos, ya que dosis muy bajas se han asociado con efectos prolongados de la droga.
- Acepromazina no posee efectos analgésicos, por lo que se deben tomar medidas adecuadas para controlar el dolor en los animales tratados.
- Usar con precaución en perros braquicéfalos (Bóxer, Pequinés, etc.) o gatos de la raza Persa, ya que pueden ser muy sensibles a los efectos bradicárdicos de la Acepromazina.
- Usar con precaución en perros con mutaciones MDR1 (Collies, Pastores australianos, entre otros), ya que pueden desarrollar una sedación más profunda y que persiste más de lo habitual.
- Razas gigantes y Galgos pueden ser extremadamente sensibles a la droga, mientras que las razas terrier son algo resistentes a sus efectos.
- Acepromazina debe usarse con precaución como agente de contención en perros agresivos, ya que puede provocar que los animales estén más propensos al sobresalto y a reaccionar a ruidos u otros estímulos sensoriales.
- Se puede utilizar Atropina en conjunto a Acepromazina para ayudar a bloquear sus efectos bradicárdicos.
- Durante la sedación con Acepromazina se recomienda utilizar lubricantes oculares o lágrimas artificiales como protectores corneales, debido a la reducción en la producción de lágrimas.
- Una dosis mayor no acelera el inicio de la sedación, solo alarga el efecto sedante.
- Una vez que se ha administrado, es importante dejar al animal en un lugar tranquilo por alrededor de 15 minutos o hasta que se logren los efectos sedantes.
- Luego de la administración, los pacientes no deben estar expuestos a fluctuaciones de temperaturas extremas por al menos 8 horas.
- La administración en conjunto con otros depresores respiratorios o del Sistema Nervioso Central, puede exacerbar la depresión respiratoria.
- Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

Adverse effects and adverse reactions:

- Accepromazine generates hypotension mediated by vasodilation, which can last for a few hours and must be considered during its use. In healthy animals this hypotension is usually mild, however it can be accentuated in anesthetized, weakened or hypovolemic patients.
- Unusual cardiovascular collapse of varying severity (secondary to bradycardia and hypotension) has been described in some dogs, especially in brachycephalic breeds.
- Accepromazine causes significant intraoperative hypothermia because it depresses the hypothalamic thermoregulatory center and causes cutaneous vasodilation. This effect lasts for a few hours.
- Accepromazine causes prolapse of the nictitating membrane, which lasts for a few hours while maintaining its effect.
- Accepromazine can decrease hematocrit for several hours, as a result of splenic congestion after α1-adrenergic receptor blockage, which may be important in anemic animals. This effect is transitory and is restored after several hours.
- Accepromazine transiently decreases platelet aggregation without affecting clotting times.
- Occasionally Accepromazine can generate contradictory clinical signs of aggressiveness and generalized stimulation of the Central Nervous System. These reactions are infrequent and transient in nature and generally do not last for more than 48 hours.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
Empty containers can be discarded as household waste, without any special precautions.
Do not dispose of containers with product residues on the ground or water courses.
For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 ° and 30° C, protected from light.

CONDITION OF SALE

Sale with Veterinary Medical prescription.

PRESENTATION

Display with 10 tablets.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

- **Chile:** Reg. SAG N° 115
- **Rep. Dominicana:** Reg. N° 8865

PANVERMIC® LARGE BREED - ORAL TABLET

COMPRIMIDO ORAL.

ANTIPARASITARIO INTERNO DE AMPLIO ESPECTRO.

Technical Specification

SPECIES

Large breed dogs.

Broad spectrum internal antiparasitic in a single dose. Effective against nematodes and cestodes, such as: *Toxocara canis, Toxascaris leonina, Ancylostoma caninum, Hookworm stenocephala, Taenia hydatigena, Taenia pisiformis, Taenia taeniformis, Dipylidium caninum, and Echinococcus granulosus.*

DOSAGE FORM

Oral Tablet

THERAPEUTIC ACTION

Broad spectrum internal antiparasitic.

COMPOSITION

Each tablet contains:
Praziquantel... ..... 175 mg
Mebendazole ... . 770 mg
Pyrantel Pamoate ... . .504 mg
(Equivalent to 175 mg of Pyrantel base)
Excipients q.s.p ..... 1 tablet

INDICATIONS

- Do not administer to dogs with liver failure, malnourished, dehydrated or anemic.
- Do not administer in dogs less than four weeks of age.
- Interactions with other pharmaceuticals:
- Do not administer together with Piperazine as it could antagonize the effect of Pyrantel Pamoate.
- Do not administer in conjunction with Morantel or Levamisole due to their mechanisms of action and toxicity similar to Pyrantel.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer orally, directly into the oral cavity or mixed with food.

- Dose of active ingredients:
Administer 22 mg / Kg of Mebendazole; 5 mg / Kg of Praziquantel and 5 mg / Kg of Pyrantel as a base.
- Product dosage:
1 tablet for every 35 kg of weight. Single dose.

For the treatment of Trichuris vulpis infestations, administer 1 tablet for every 35 kg of weight for 3 consecutive days.

CONTRAINDICATIONS

- Do not administer to dogs with liver failure, malnourished, dehydrated or anemic.
- Do not administer in dogs less than four weeks of age.
- Interactions with other pharmaceuticals:
- Do not administer together with Piperazine as it could antagonize the effect of Pyrantel Pamoate.
- Do not administer in conjunction with Morantel or Levamisole due to their mechanisms of action and toxicity similar to Pyrantel.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Wash your hands after administering the product.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

- In some animals loose stools and in some cases moderate diarrhea may occur.
- In less than 5% of treated dogs there may be anorexia, vomiting and lethargy.

CONSERVATION

- Keep in a dry place and protected from light.
- Store at room temperature between 15 and 30 ° C

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

Box with 1 tablets

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 2076
Bolivia: Reg. SENASAG PUV-F-N° 005508/13

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

PANVERMIC® ORAL TABLET

COMPRIMIDO ORAL.

ANTIPARASITARIO INTERNO DE AMPLIO ESPECTRO.

Technical Specification

SPECIES

Dogs.

Broad spectrum internal antiparasitic in a single dose. Effective against nematodes and cestodes, such as: *Toxocara canis, Toxascaris leonina, Ancylostoma caninum, Hookworm stenocephala, Taenia hydatigena, Taenia pisiformis, Taenia taeniformis, Dipylidium caninum, and Echinococcus granulosus.*

DOSAGE FORM

Oral Tablet

THERAPEUTIC ACTION

Broad spectrum internal antiparasitic.

COMPOSITION

Each tablet contains:
Praziquantel ..... 50 mg
Mebendazole ..... 220 mg
Pyrantel Pamoate ..... 144 mg
(Equivalent to 50 mg of Pyrantel base)
Excipients c.s.p ..... 1 tablet

INDICATIONS

- Do not administer in dogs with known hypersensitivity to the active ingredients.
- Do not administer to puppies under 4 weeks of age.
- Do not administer to dogs with liver failure, malnourished, dehydrated or anemic.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer orally, directly into the oral cavity or mixed with food.

- Dose of active ingredients:
Administer 22 mg / Kg of Mebendazole; 5 mg / Kg of Praziquantel and 5 mg / Kg of Pyrantel as a base.
- Product dosage:
1 tablet for every 10 kg of weight. Single dose.

For the treatment of Trichuris vulpis infestations, administer for 3 consecutive days.

DRUG INTERACTIONS

- Do not administer together with Piperazine as it could antagonize the effect of Pyrantel Pamoate.
- Do not administer in conjunction with Morantel or Levamisole due to their mechanisms of action and toxicity similar to Pyrantel.

CONTRAINDICATIONS

- Do not administer in dogs with known hypersensitivity to the active ingredients.
- Do not administer to puppies under 4 weeks of age.
- Do not administer to dogs with liver failure, malnourished, dehydrated or anemic.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Wash your hands after administering the product.
- In case of accidental ingestion, do not induce vomiting. Get medical help.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

In some animals they can present in a mild and transient way: nausea, vomiting, anorexia, abdominal pain and diarrhea.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of natural water. Dispose of this product waste carefully with household waste.

CONSERVATION

Keep in a cool and dry place, at room temperature between 15° and 30° C.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

Box with 2 tablets

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Reg. SAG N° 1689

PAPAINPET® ORAL TABLET.

COMPRIMIDO ORAL.

SUPLEMENTO NUTRICIONAL PARA PERROS Y GATOS.

Technical Specification

SPECIES

Dogs and cats

Papainpet® is a nutritional supplement based on papaya extract specially formulated for dogs and cats. Papaya extract is a supplement rich in papain, a proteolytic enzyme with various beneficial properties for your pet.

DOSAGE FORM

Oral tablet.

THERAPEUTIC ACTION

Nutritional supplement for dogs and cats.

COMPOSITION

Each tablet contains:
Papaya Extract .....50 mg
Excipients q.s.p .....1 tablet

ROUTE OD ADMINISTRATION AND DOSAGE

Route of administration: Oral route.

Suggested dosage:
- Dogs: 1 tablet every 10 kg of weight, 2 or 3 times a day.
- Cats: ½ tablet every 4 Kg of weight, 1 or 2 times a day.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

- EXCLUSIVE USE IN ANIMAL FEED
- DOES NOT CORRESPOND TO A COMPLETE FOOD

CONSERVATION

Keep in a cool, dry place, protected from light, at no more than 30° C.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

30 tablets

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Reg. LENAA N°: RM 03-008N

PAZ-PET® ORAL SUSPENSION

SUSPENSIÓN ORAL.

MODIFICADOR DE CONDUCTA PARA PERROS.

Technical Specification

SPECIES

Dogs.

- PAZ-PET® is very useful for your pet when it has to face very stressful situations, such as: Environmental changes, transfers.
- Excessive barking or vocalization Accommodation in canine hotels or similar facilities.
- Anxiety states over separation from its owners.
- Socialization with other animals.
- Noisy or overstimulating environments for dog (parties, fireworks, nearby airports, etc.)
- Behavior problems such as attempts to mate, excessive walking, self-induced injuries commonly observed in dogs confined in small spaces.
- Do not cause sleepiness or depression.

DOSAGE FORM

Oral suspension.

THERAPEUTIC ACTION

Behavior modifier for dogs.

COMPOSITION

Each mL contains:

L-Theanine.....30 mg

Thiamine (vitamin B1).....30 mg

L-Tryptophan.....3 mg

Vegetal concentrate.....30 mg

(*Passiflora sp., Valeriana sp., Melissa sp.*)

Excipients q.s.p.....1 mL

USE INSTRUCTIONS

Shake before use.

ROUTE OD ADMINISTRATION AND DOSAGE

2 mL each 10 kg of body weight, once daily. Oral route. The effect onset occurs approximately at the third day of the dosage beginning. It is recommended to administer the product for at least 30 days.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a fresh, dry place, out of direct light, at no more than 30°C.

PRESENTATION

60 mL

PREPARED BY

Drag Pharma Laboratory.

PETEVER® FORTE - SHAMPOO

SHAMPOO

SHAMPOO ANTISÉPTICO

Technical Specification

SPECIES

Dogs.

Petever® shampoo, due to its effective antiseptic action against skin bacteria and yeasts, represents an effective aid in the treatment of superficial and deep skin infections caused by *Staphylococcus intermedius* and *Malassezia pachydermatis* in dogs.

DOSAGE FORM

Shampoo

THERAPEUTIC ACTION

Antiseptic shampoo

COMPOSITION

Each 100 mL contains:
Chlorhexidine Gluconate ..... 3 g
Excipients q.s.p ..... 100 mL

MODE OF APPLICATION

- Moisten the animal with lukewarm water.
- Apply a sufficient amount of the shampoo to several points, massage and rinse.
- Apply again, massage and leave for 10 minutes. Then rinse with plenty of clean water.
- Repeat two or three times a week until the symptoms subside and then once a week or as indicated by the Veterinarian.

ROUTE OD ADMINISTRATION AND DOSAGE

Topical Use

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a cool and dry place, at room temperature between 15 and 30 ° C.
Avoid the direct exposure to the sun.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

150 mL bottle

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 1921
Uruguay: Reg. MGAP: 2017A00501
Perú: Reg. SENASA F.G6.21.I.0003

COUNTRIES WHERE IT IS MARKETED

Importer in Uruguay by:
VIVAFIL S.A. RIO NEGRO 1107
Montevideo - Uruguay, TEL 29001112
grupotecnovet@gmail.com
Technical Director: DMTV Diego Cuadrado.

Imported and Distributed in Peru by Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

PETEVER® PLUS - ORAL SOLUTION

SOLUCIÓN ORAL.

ANTISÉPTICO BUCAL.

Technical Specification

SPECIES

Dogs

Petever® Plus, Oral solution, is indicated as an aid for the maintenance of oral health and the control of halitosis in dogs. Its use in combination with daily tooth brushing helps remove food particles, preventing the deposit of bacterial plaque, dental stones and the appearance of gingivitis. To maximize the results in reducing gingivitis and plaque deposition, it is recommended to be used after dental scaling and polishing.

DOSAGE FORM

Oral Solution

THERAPEUTIC ACTION

Oral antiseptic.

COMPOSITION

Each 100 mL of oral solution contains:
Chlorhexidine gluconate ..... 0.12 g
Ethyl Alcohol ..... 6.00 g
Excipients q.s.p ..... 100 mL

INDICATIONS

- It is recommended not to administer in individuals with known hypersensitivity to chlorhexidine or any component of the preparation.
- Avoid its administration in conjunction with common detergents or anionic compounds, since they inactivate the action of Chlorhexidine.

MODE OF APPLICATION

- **As a mouthwash:** Support the animal's head with one hand, and spray the indicated dose of Petever® Plus, Oral Solution, on the surface of the teeth and gums on one side of the animal's snout. Massage the animal's snout on the teeth so that the solution acts on the surface of the teeth. Then repeat the operation on the other side of the muzzle.
- **Along with tooth brushing:** Place a small amount (about 3 teaspoons) of Petever® Plus, Oral Solution, in a clean container or glass. Support the animal's head with one hand, and dip the toothbrush in the solution and carefully brush your pet's teeth in a gentle motion. Dip the brush in the solution as many times as necessary.

ROUTE OD ADMINISTRATION AND DOSAGE

Topical oral administration in atomized form.

Recommended dose:
Apply 3 sprays to each tooth, once a day, as a mouthwash or along with your pet's regular tooth brushing.

CONTRAINDICATIONS

- It is recommended not to administer in individuals with known hypersensitivity to chlorhexidine or any component of the preparation.
- Avoid its administration in conjunction with common detergents or anionic compounds, since they inactivate the action of Chlorhexidine.

PRECAUTIONS

- The use of PETEVER® PLUS, oral solution, is not a substitute for veterinary dental treatment
- This product should be used under the supervision of a Veterinarian who suggests a complete oral health program for your pet.
- Keep out of the reach of children.
- This product can be administered to pregnant or lactating females.

SPECIAL PRECAUTIONS FOR USE

- The use of PETEVER® PLUS, oral solution, is not a substitute for veterinary dental treatment
- This product should be used under the supervision of a Veterinarian who suggests a complete oral health program for your pet.
- Keep out of the reach of children.
- This product can be administered to pregnant or lactating females.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Flammable product, keep away from heat sources.

SIDE EFFECTS

Use for periods longer than 30 days can cause unwanted stains on tooth surfaces.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:

- Dispose of unused product remains in its original container, well closed.
- Dispose of the waste of this product with care together with household waste.

CONSERVATION

- Store in a cool place (between 15 ° and 30 ° C), dry and protected from light.
- Keep away from heat sources, flammable product.

CONDITION OF SALE

Free sale.

PRESENTATION

100 mL spray bottle

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 1932
Perú: Registro SENASA F.58.32.I.0084

PET-OTIC® OTIC SOLUTION

SOLUCIÓN ÓTICA.

SOLUCIÓN DE LIMPIEZA ÓTICA.

Technical Specification

SPECIES

Dogs and cats.

Pet-Otic® is an effective combination of ceruminolytic agents and emulsifiers specially formulated to clean the ears of dogs and cats. It removes the wax and reduces the bad smell of the ear.
Pet-Otic® removes cellular debris and necrotic debris from superficial wet skin lesions. The presence of organic acids in its composition gives it an effective drying and acidifying action, which favors the removal of cell debris and provides an optimal environment for cell regeneration.
Pet-Otic® can be used in routine grooming of your pet's ear. Does not contain alcohol.

DOSAGE FORM

Otic Solution

THERAPEUTIC ACTION

Otic cleaning solution

COMPOSITION

- Acetic Acid 2%
- Boric acid 2%
- Eucalyptus oil, Aloe Vera and surfactants.

MODE OF APPLICATION

- Place the upper extremity of Pet-Otic® in the ear and press to freely apply the solution in the ear canal area.
- Gently massage from the base of the ear to the top of the ear. Use 2 or 3 times a day until symptoms stop.
- For routine cleaning, carefully administer Pet-OticC® in the ear and massage at the base of the ear to loosen any dirt.
- Use a cotton ball to remove any residue and excess solution.
- Apply 2-3 times a week.

ROUTE OD ADMINISTRATION AND DOSAGE

Otic administration.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

store in a cool, dry place and away from light, at no more than 30ºC.

CONDITION OF SALE

OTC non prescription

PRESENTATION

Dropper bottle with 100 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

PET OUT® SPRAY

SPRAY.

REPELENTE PARA PERROS Y GATOS.

Technical Specification

SPECIES

Dogs and cats.

Pet Out® is a product specially formulated for pushing dogs and cats away from sites or objects determined by the owner. Pet Out® helps to end with the problems caused by the territory marking and bad toilet behavior of dogs and cats. Pet Out® is applied in the form of spray on the surface chosen. This must be repeated several times daily during the first days, until your pet learns that the use of such a place is restricted. Pet Out® must not be applied directly on the coat of your pet; if this occurs, wash with plenty of water and rinse repeatedly. Pet Out® must not be applied directly on your pet's eyes.

DOSAGE FORM

Spray.

THERAPEUTIC ACTION

Dogs and cats repellent.

INGREDIENTS

Citronella Oil, Isopropyl Alcohol, 96% Denatured Ethyl Alcohol, Nonyl Methyl Ketone, Purified Water.

WARNINGS

- Mantener fuera del alcance de los niños.
- Inflamable.

Advertencias al usuario:
Los patrones de comportamiento y de marcaje de las mascotas están regulados por numerosos factores. Este producto pudiese no interrumpir los patrones de micción y/o defecación establecidos en perros y gatos, a menos que se utilice como una herramienta complementaria de entrenamiento.

CONSERVATION

Store in a cool, dry place, protected from light, below 30 °C.

PRESENTATION

spray with 160 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Costa Rica: Reg. MAG N° CL4-13-2-6021

PICHICHI® SOLUTION

SOLUCIÓN.

EDUCADOR SANITARIO PARA PERROS.

Technical Specification

SPECIES

Dogs.

Potty trainer for dogs. Teach your dog where to pee and poop.

DOSAGE FORM

Solution.

THERAPEUTIC ACTION

Potty trainer for dogs.

COMPOSITION

Demineralized water, Ammonium chloride, Sodium bicarbonate, Ammonia, Colorant.

MODE OF APPLICATION

Apply 5 - 10 drops of PICHICHI® in the place where you want your dog pees and poops.

Make sure your pet smells the place where you applied PICHICHI® in order to get an optimal result.

Apply it directly on a cloth, cardboard o newspaper and let it in the intended place since they preserve the product action longer. Repeat frequently this operation, especially after meals, at the morning and evening.

Apply PICHICHI® every day until your pet becomes accustomed to pee and poop in the intended place; then, it is only necessary to apply PICHICHI® once or twice a week.

ROUTE OD ADMINISTRATION AND DOSAGE

For external use.

WARNINGS

- Este producto no debe ser ingerido.
- Mantener fuera del alcance de los niños.

PRESENTATION

20 mL

PREPARED BY

Drag Pharma Laboratory.

PRAZIVERMIC® ORAL TABLET

COMPRIMIDO ORAL.

ANTIPARASITARIO INTERNO DE AMPLIO ESPECTRO.

Technical Specification

SPECIES

Dogs.

Broad spectrum anthelmintic, effective against cestodes and nematodes present in dogs, such as: *Toxocara canis, Ancylostoma caninum, Taenia hydatigena, Taenia pisiformis, Echinococcus granulosus, and Dipylidium caninum.*

DOSAGE FORM

Oral tablet

THERAPEUTIC ACTION

Broad spectrum internal antiparasitic.

COMPOSITION

Each tablet contains:
Levamisole Hydrochloride ..... 80 mg
Praziquantel ..... 50 mg
Excipients q.s.p ....,,,,..... 1 tab.

INDICATIONS

- Do not administer to dogs under 4 weeks of age.
- Do not administer to patients with a history of gastric intolerance.
- Do not administer to animals with hypersensitivity to any of the active principles.

Use during pregnancy and lactation:
Do not administer to pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Route of administration and method of use: Oral route.
Administer directly into the oral cavity or mixed with food.

Dose:
Administer 1 tablet for every 10 kilos of weight, equivalent to 8 mg / Kg of Levamisole Hydrochloride and 5 mg / Kg of Praziquantel, in a single dose.

DRUG INTERACTIONS

- It is recommended to avoid the administration of Levamisole in conjunction with Chloramphenicol, since deaths have been reported when they have been administered concomitantly.
- Cholinesterase inhibitor drugs (eg, organophosphates, Neostigmine) can potentiate the toxic effects of Levamisole.
- Nicotinic-type components (eg, Pyrantel, Morantel, Diethylcarbamazine) can enhance the toxic effects of Levamisole.
- Levamisole administered together with Warfarin increases the risk of bleeding.
- Interactions of Praziquantel with other medicinal products for veterinary use have not been described.

CONTRAINDICATIONS

- Do not administer to dogs under 4 weeks of age.
- Do not administer to patients with a history of gastric intolerance.
- Do not administer to animals with hypersensitivity to any of the active principles.

Use during pregnancy and lactation:
Do not administer to pregnant or lactating females.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Wash your hands with soap and plenty of water after handling the product.
- In the case of contact with the eyes, it is recommended to wash with plenty of water for 15 minutes. If irritation develops and persists, see doctor.
- In case of accidental ingestion, do not induce vomiting. Get medical help.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

In some cases, hypersalivation, vomiting or muscle tremors may be observed, which are temporary symptoms and disappear spontaneously.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of natural water. Do not reuse the container. Dispose of this product waste carefully with household waste.

CONSERVATION

Store in a cool, dry place, at room temperature between 15° and 30° C.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

Box with 2 tablets

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 0362
Perú: Registro SENASA F.08.21.I.0925

COUNTRIES WHERE IT IS MARKETED

Imported and Distributed in Peru by:
Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

QUIT OLOR® SPRAY SOLUTION

SPRAY.

Technical Specification

SPECIES

Dogs and cats.

QUIT OLOR® is indicated to eliminate unpleasant odors from organic waste of any pet, such as feces, urine, vomiting, exudates, territory marker, among others. It may be applied on tapestries, carpets, curtains, car's interior, terraces, corners, rooms, warehouses, environments, etc.

DOSAGE FORM

Spray solution.

INGREDIENTS

Purified water, Odoriferous 23DP, Propylenglycol, Ethanol, Cetyl Trimethyl Ammonium Chloride, Cocamide DEA, Nonylphenol ethoxylate, authorized flavorings.

PROPERTIES

Odor neutralizer.

USE INSTRUCTIONS

- For environments, spray QUIT OLOR® covering the most part of the intended space.
- For surfaces, apply a quantity enough to moisten the surface from where the unpleasant odor arises.
- Repeat the procedure 2 to 4 times daily, as your own evaluation.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

QUIT OLOR® does not stain or change the color of fabrics, rugs, carpets or curtains.

PRESENTATION

500 mL

PREPARED BY

Drag Pharma Laboratory.

REGEPIPEL® PLUS - SHAMPOO

SHAMPOO.

ANTISÉPTICO Y ANTIMICÓTICO.

Technical Specification

SPECIES

Dogs and cats.

For the treatment of seborrheic dermatitis associated with *Staphylococcus intermedius* infections, *Malassezia pachydermatis*, and as an aid in the treatment of dermatophyte infections due to *Microsporum canis* in dogs and cats.

DOSAGE FORM

Shampoo.

THERAPEUTIC ACTION

Antifungal and antiseptic shampoo.

COMPOSITION

Each 100 mL contains:
Chlorhexidine Gluconate 20% ..... 10 g
(Equivalent to 2 g of Chlorhexidine Gluconate)
Miconazole Nitrate ..... 2.3 g
(Equivalent to 2 g of Miconazole base)
Excipients q.s.p ..... 100 mL

INDICATIONS

- Do not use in cats younger than 6 weeks of age and in dogs younger than 4 weeks of age.
- Do not administer to individuals with known hypersensitivity to any component of the preparation.
- Do not use in pregnant or lactating females.

MODE OF APPLICATION

- Wet the animal with plenty of water.
- Apply a sufficient amount of the shampoo at various points and massage. With the help of a sponge, spread to promote its penetration into the skin and coat. Be sure to apply Regepipel® Plus around the lips, under the tail and between the fingers as these are reservoir areas for causative agents.
- Leave on for 10 to 15 minutes, then rinse with plenty of clean water.
- Repeat two or three times a week until symptoms subside and then once a week or as directed by the Veterinary Doctor.

ROUTE OD ADMINISTRATION AND DOSAGE

Topically.

DRUG INTERACTIONS

Chlorhexidine is inactivated when used with common detergents or anionic compounds.

CONTRAINDICATIONS

- Do not use in cats younger than 6 weeks of age and in dogs younger than 4 weeks of age.
- Do not administer to individuals with known hypersensitivity to any component of the preparation.
- Do not use in pregnant or lactating females.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Avoid contact with eyes, in case of accidental contact wash immediately with plenty of water.

WARNINGS

Advertencias y precauciones especiales de uso:

- En el tratamiento de dermatofitosis, la asociación de la terapia tópica a la terapia sistémica favorece la cura de las lesiones en 6-9 semanas, tiempo menor al requerido cuando se administra sólo la terapia sistémica.
- En caso de irritación local, suspender el tratamiento.
- Evitar el contacto con los ojos del animal, en caso de contacto accidental lavar inmediatamente con abundante agua.
- Mantener fuera del alcance de los niños.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
Empty containers can be discarded as household waste, without any special precautions. Do not dispose of containers with product residues on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Keep at room temperature, between 15 and 30°C.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Bottle with 150 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 1696
Panamá: Reg. N° RF-8355-18
El Salvador: VE2013084783
Bolivia: Reg. SENASAG PUV-F N° 007259/16
Uruguay: Reg. MGAP N° 2019A00682

COUNTRIES WHERE IT IS MARKETED

Distribution in El Salvador: Rafael Alfredo Alfaro Castillo.
8a C. Pte. Pje. Moreno N ° 112, Col. Flor Blanca.
San Salvador, El Salvador.

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

Imported in Uruguay by: VIVAFIL S.A.
RIO NEGRO 1107 Montevideo - Uruguay, TEL 29001112
grupotecnovet@gmail.com
Technical Director: DMTV Diego Cuadrado.

RESPIG® ORAL SUSPENSION

SUSPENSIÓN ORAL.

ANTIMICROBIANO DE AMPLIO ESPECTRO.

Technical Specification

SPECIES

Dogs and cats.

Broad spectrum antimicrobial indicated for the treatment of respiratory tract and urinary tract infections and soft tissue infections caused by Gram positive and Gram negative bacteria susceptible to Sulfamethoxazole and Trimethoprim.

DOSAGE FORM

Oral suspension

THERAPEUTIC ACTION

Broad spectrum antimicrobial

COMPOSITION

Each 5 mL of oral suspension contains:
Sulfamethoxazole ..... 200 mg
Trimethoprim ..... 40 mg
Excipients q.s. ..... 5 mL

INDICATIONS

- Do not use in dogs with liver and / or kidney damage.
- Do not use in dogs with hypersensitivity to sulfonamides.
- Do not use in Pinscher breed dogs.
- Do not use in pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Orally.

- **Dosage of active ingredients:**
  30 mg of the association per kilo of weight every 24 hours, or 15 mg of the association per kilo of weight every 12 hours, for 7 to 10 days.
- **Product dosage:**
  5 mL for every 8 Kg of weight, every 24 hours; or 2.5 mL every 8 Kg of weight, every 12 hours, for 7 to 10 days.

Do not administer for more than 14 days.

DRUG INTERACTIONS

- The association of Sulfamethoxazole and Trimethoprim may prolong clotting times in patients medicated with coumarin anticoagulants.
- Antacids can decrease the bioavailability of sulfonamides if administered concurrently.
- Trimethoprim can reduce the therapeutic effects of cyclosporine (systemic) and increase the risk of nephrotoxicity.

CONTRAINDICATIONS

- Do not use in dogs with liver and / or kidney damage.
- Do not use in dogs with hypersensitivity to sulfonamides.
- Do not use in Pinscher breed dogs.
- Do not use in pregnant or lactating females.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Do not handle by people hypersensitive to sulfonamides.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 ° C, protected from light.
Once the container is opened, use within 14 days. Discard the unused product after that period of time

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

100 mL bottle

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 944

RHINOLIN® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

MUCOLÍTICO.

Technical Specification

SPECIES

Horses, cattle, dogs and cats.

Indicated in the treatment of respiratory diseases associated with increased production or an increase in the viscosity of the bronchial mucus, such as acute or chronic bronchitis, acute or chronic bronchopneumonia, rhinitis. It is also indicated as support therapy for specific antimicrobial treatment.

DOSAGE FORM

Injectable solution

THERAPEUTIC ACTION

Mucolytic

COMPOSITION

Each 1 mL contains:
Bromhexine Hydrochloride ..... 3 mg
Excipients q.s. ..... 1 mL

INDICATIONS

- Do not administer in cases of pulmonary edema.
- Do not administer in animals with known hypersensitivity to Bromhexine.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration by intramuscular route.

Dose of the active principle:

- **Horses:** 0.15 mg / Kg.
- **Cattle, dogs and cats:** 0.5 mg / Kg.

Product dosage:

- **Horses:** 5 mL / 100 kilos.
  (As a reference, foals: 10 to 15 mL / day; adult horses: 20 to 30 mL / day).
- **Cattle:** 5 mL / 30 kilos.
  (As reference calves: 5 to 30 mL / day; adult bovines: 30 to 65 mL / day).
- **Dogs**: 2.5 mL / 15 kilos.
- **Cats:** 0.5 mL / 3 kilos.

Frequency and duration of treatment:
Administer every 24 hours. To achieve an optimal effect, administration for at least 5 days according to the treating Veterinarian is recommended.

CONTRAINDICATIONS

- Do not administer in cases of pulmonary edema.
- Do not administer in animals with known hypersensitivity to Bromhexine.

PRECAUTIONS

- Keep out of the reach of children.
- If the required dose is greater than 20 mL, it is suggested to divide it into two or more application points.
- In the case of an infectious etiology, it must be accompanied in its use with a specific antibiotic.

SPECIAL PRECAUTIONS FOR USE

- Keep out of the reach of children.
- If the required dose is greater than 20 mL, it is suggested to divide it into two or more application points.
- In the case of an infectious etiology, it must be accompanied in its use with a specific antibiotic.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- In the case of ocular exposure, irrigate the eyes with water at room temperature for at least 15 minutes. If irritation exists or persists, consult a doctor.
- In the case of dermal exposure, wash the area with soap and water. If irritation exists or persists, consult a doctor.
- In the case of accidental injection, medical attention should be sought.

WARNINGS

GUARD PERIOD

- **Beef cattle:** 0 days.
  Do not administer to bovines whose milk is intended for human consumption.
  Do not administer in horses whose meat is intended for human consumption.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15° and 30°C, protected from light.
Once the container is opened, use within 28 days. Discard unused product after that time frame.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

100 mL vial

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

- **Chile:** Reg. SAG N° 1815
- **Rep. Dominicana:** Reg. Nº 6343
- **Panamá:** Reg. N° RF-8354-18

ROSTRUM® 10% - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANTIBACTERIANO DE AMPIO ESPECTRO.

Technical Specification

SPECIES

Pigs, Bovines and Sheep.

Broad spectrum antibacterial recommended for the treatment and prophylaxis of infectious diseases caused by bacteria sensitive to Enrofloxacin. Rostrum® is effective against Gram negative and Gram positive bacteria, such as: *E. coli, Salmonella spp., Pasteurella spp., Haemophilus spp., Bordetella spp., Campylobacter spp., Proteus spp., Klebsiella spp., Erysipelothrix spp. , Streptococcus spp., Staphylococcus spp. and Mycoplasma spp.*

Rostrum® is recommended for the treatment of infections of the gastrointestinal, respiratory, genito-urinary and dermal tracts and in cases of contaminated wounds.

- Bovines: Diarrhea, Pneumonia, Salmonellosis, Mastitis, Metritis, Omphalitis, Septicemia and secondary infections in viral process.

- Sheep: diarrhea, pneumonia, acute mastitis, gangrenous mastitis, contagious agalactia.

- Pigs: Pre and post weaning diarrhea, Pleuropneumonia (*Haemophilus spp.*), Salmonellosis, Ileitis, Atrophic rhinitis, Epidermitis, Erysipelas, Enzootic pneumonia (*Mycoplasma spp.*), Mastitis syndrome, Metritis, Agalactia (M.M.A)

DOSAGE FORM

Injectable Solution

THERAPEUTIC ACTION

Broad spectrum antibacterial

COMPOSITION

Each 100 mL of solution for injection contains:
Enrofloxacin ..... 10 g
Excipients q.s.p ..... 100 mL

INDICATIONS

- Do not administer Enrofloxacin to animals with hypersensitivity to quinolones.
- The use of Enrofloxacin can cause the formation of crystals in the urine, for which it is recommended not to administer to dehydrated animals.
- Do not administer in pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Parenteral route.

Intramuscular in pigs.
Subcutaneous in cattle and sheep.

Product dosage:

- **Cattle, sheep and pigs** 1 mL / 40 kg of weight (equivalent to 2.5 mg / kg of weight), every 24 hours for 3 days. In the case of complicated respiratory infections and Salmonellosis, it is recommended to increase the dose to 1 mL / 20 kg of weight (equivalent to 5 mg / Kg of weight) and for 5 days in the case of Salmonellosis.
- **In sheep**, a dose of 1 mL / 20 kg of weight (equivalent to 5 mg / kg of weight) is recommended in the case of gangrenous mastitis, acute mastitis and contagious agalactia, for 3 days.

DRUG INTERACTIONS

Enrofloxacin administered with theophylline can increase your serum levels.
Concomitant administration with other drugs that have hepatic metabolism may affect the pharmacokinetics of one or both drugs.

CONTRAINDICATIONS

- Do not administer Enrofloxacin to animals with hypersensitivity to quinolones.
- The use of Enrofloxacin can cause the formation of crystals in the urine, for which it is recommended not to administer to dehydrated animals.
- Do not administer in pregnant or lactating females.

PRECAUTIONS

- Avoid the use of Enrofloxacin in young growing animals, due to the cartilaginous abnormalities that it can generate.
- Fluoroquinolones, like Enrofloxacin, should not be used as the first line of treatment, unless there is no therapeutic alternative available. When used as a second treatment, it should be based on susceptibility studies.

SPECIAL PRECAUTIONS FOR USE

- Avoid the use of Enrofloxacin in young growing animals, due to the cartilaginous abnormalities that it can generate.
- Fluoroquinolones, like Enrofloxacin, should not be used as the first line of treatment, unless there is no therapeutic alternative available. When used as a second treatment, it should be based on susceptibility studies.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Wash your hands after administering the product.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

Arthropathies have been reported in immature animals (calves, piglets). The risk of arthropathy increases when the dose is increased, however, it has also been reported when the recommended doses are used.

Other adverse effects that have been reported in very rare cases (less than 1 animal in 10,000 animals, including isolated reports) include vomiting, anorexia, increased liver enzymes, ataxia, seizures, depression, lethargy and nervousness.

GUARD PERIOD

Meat:

- Cattle and sheep: 14 days.
- Pigs: 10 days.

Milk: Do not administer in animals whose milk is intended for human consumption.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of water.
Discard the empty container or with product remains together with household waste.

CONSERVATION

Store between 15 and 30 °C, protected from light.
Once the container is opened, use within 28 days. Discard unused product after that time frame.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

100 mL vial

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 600
El Salvador: VE2006053414

COUNTRIES WHERE IT IS MARKETED

Dist. In El Salvador by: Rafael Alfredo Alfaro Castillo.

ROSTRUM® 2.5% - ORAL SUSPENSION

SUSPENSIÓN ORAL.

ANTIBACTERIANO.

Technical Specification

SPECIES

Dogs and cats.

Rostrum®, is a broad spectrum antibacterial, recommended for the treatment of:

- Gastrointestinal tract infections (*Escherichia coli, Salmonella spp., Proteus spp.*).
- Respiratory tract infections (*Pasteurella spp., Bordetella spp., Klebsiella spp.*).
- Genitourinary infections such as nephritis, pyelonephritis, cystitis. (*Escherichia coli, Corynebacteriumpyogenes, Staphylococcus aureus, Streptococcus spp. And Pseudomonas aeruginosa*).
- Skin infections
- Infections of the external auditory canal and otitis (*Escherichia coli, Staphylococcus aureus, Streptococcus spp. And Pseudomonas aeruginosa*).
- Dermatitis, infected wounds (*Escherichia coli, Klebsiella spp., Staphylococcus aureus, Streptococcus spp. And Pseudomonas aeruginosa*).

DOSAGE FORM

Oral Suspension

THERAPEUTIC ACTION

Antibacterial

COMPOSITION

Each mL of oral suspension contains:
Enrofloxacin ..... 25 mg
Excipients c.s.p. ..... 1 mL

PROPERTIES

Enrofloxacin is a synthetic antimicrobial agent belonging to the Fluoroquinolone family, which are characterized by being bactericidal antibiotics that have powerful antibacterial activity against a wide variety of microorganisms, mainly against Gram negative, Gram positive, Mycoplasma and Chlamydial bacteria.

INDICATIONS

- Enrofloxacin is contraindicated in animals with hypersensitivity to quinolones.
- Do not administer to small to medium breed dogs under 1 year old, or to large breed dogs under 1 ½ years.
- Do not administer to cats under 8 weeks of age.
- The use of Enrofloxacin can generate the formation of crystals in urine, for which it is recommended not to administer in dehydrated animals.
- Do not administer in pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Via oral administration.

Dose of the active principle:

- **Dogs:** 5 - 20 mg / Kg of body weight once a day for 7-10 days. The dose can be administered as a single dose or divided into two equal doses, administered every 12 hours.
- **Cats:** 5 mg / Kg of body weight once a day for 7-10 days. The dose can be administered as a single dose or divided into two equal doses, administered every 12 hours. In case of mycoplasma infection, the duration of treatment should be 14 to 21 days.

Product dosage:

- **Dogs:** 1 to 4 mL every 5 Kg of body weight every 24 hours * for 7 - 10 days.
- **Cats:** 1 mL every 5 Kg of body weight every 24 hours * for 7-10 days.

* The dose can be administered as a single dose or divided into two equal doses, administered every 12 hours. In case of mycoplasmosis, the duration of treatment should be 14 to 21 days.

DRUG INTERACTIONS

- Compounds containing di or trivalent cations such as Aluminum, Calcium, Magnesium, Iron or Zinc can decrease the absorption of Enrofloxacin. Thus, for example antacids containing cations (Mg2 +, Al3 +, Ca2 +) can bind Enrofloxacin and prevent its absorption.
- Sucralfate can inhibit the absorption of Enrofloxacin. It is recommended to separate both dosages for at least two hours.
- Enrofloxacin administered with Theophylline can increase your serum levels. In dogs, theophylline clearance was reduced by 43% when it was administered in conjunction with Enrofloxacin at a dose of 5 mg / Kg every 24 hours.
- Concomitant administration with other drugs that have hepatic metabolism may affect the pharmacokinetics of one or both drugs.

CONTRAINDICATIONS

- Enrofloxacin is contraindicated in animals with hypersensitivity to quinolones.
- Do not administer to small to medium breed dogs under 1 year old, or to large breed dogs under 1 ½ years.
- Do not administer to cats under 8 weeks of age.
- The use of Enrofloxacin can generate the formation of crystals in urine, for which it is recommended not to administer in dehydrated animals.
- Do not administer in pregnant or lactating females.

PRECAUTIONS

Fluoroquinolones, such as Enrofloxacin, should not be used as the first line of treatment, unless there is no therapeutic alternative available. When used as a second treatment, it should be based on susceptibility studies.

SPECIAL PRECAUTIONS FOR USE

Fluoroquinolones, such as Enrofloxacin, should not be used as the first line of treatment, unless there is no therapeutic alternative available. When used as a second treatment, it should be based on susceptibility studies.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Wash your hands after administering the product.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

- Arthropathies have been reported in immature animals, especially dogs. The risk of arthropathy increases when the dose is increased, however it has also been reported when the recommended doses in dogs are used.
- Other adverse effects that have been reported in very rare cases (less than 1 animal in 10,000 animals, including isolated reports) include vomiting, anorexia, increased liver enzymes, ataxia, seizures, depression, lethargy and nervousness.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of natural water. Dispose of this product waste carefully with household waste.

CONSERVATION

- Store at room temperature between 15 and 30 ° C in a dry place.
- Protect yourself from light.
- Do not freeze.
- Once opened, use the product within 3 months. Discard the unused product after that period of time.

CONDITION OF SALE

Sale with retained Veterinary Medical prescription.

PRESENTATION

20 mL bottle

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. S.A.G. N° 2038
Bolivia: Reg. SENASAG PUV-F-N° 006146/14
Costa Rica: MAG CL4-5-60-6094

ROSTRUM® 50 MG - ORAL TABLET

COMPRIMIDO ORAL.

ANTIBACTERIANO.

Technical Specification

SPECIES

Dogs and cats.

Rostrum®, is a broad spectrum antibacterial, recommended for the treatment of:

- Gastrointestinal tract infections (*Escherichia coli, Salmonella spp., Proteus spp.*).
- Respiratory tract infections (*Pasteurella spp., Bordetella spp., Klebsiella spp.*).
- Genitourinary infections such as nephritis, pyelonephritis, cystitis. (*Escherichia coli, Corynebacterium pyogenes, Staphylococcus aureus, Streptococcus spp. And Pseudomonas aeruginosa*).
- Skin infections
- External auditory canal infections and otitis (*Escherichia coli, Staphylococcus aureus, Streptococcus spp. And Pseudomonas aeruginosa*).
- Dermatitis, infected wounds (*Escherichia coli, Klebsiella spp., Staphylococcus aureus, Streptococcus spp. And Pseudomonas aeruginosa*).

DOSAGE FORM

Oral Tablet

THERAPEUTIC ACTION

Antibacterial

COMPOSITION

Each tablet contains:
Enrofloxacin ..... 50 mg
Excipients q.s.p ..... 1 tablet

INDICATIONS

- Enrofloxacin is contraindicated in animals with hypersensitivity to quinolones.
- Do not administer to small to medium breed dogs, younger than 1 year, nor to large breed dogs younger than 1 ½ year.
- Do not administer to cats younger than 8 weeks of age.
- The use of Enrofloxacin can cause the formation of crystals in urine, therefore it is recommended not to administer in dehydrated animals.
- It is not recommended for use in pregnancy and / or lactation.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration.

Dosage: 1 tablet of 50 mg / 10 Kg of weight per day, for 7 to 10 days, equivalent to 5 mg / Kg of weight / day / for 7 to 10 days.

Half the dose can be administered twice a day.

DRUG INTERACTIONS

- Compounds containing di or trivalent cations such as aluminum, calcium, magnesium, iron or zinc can decrease the absorption of Enrofloxacin. Thus, for example, antacids containing cations (Mg2 +, Al3 +, Ca2 +) can bind to Enrofloxacin and prevent its absorption.
- Sucralfate can inhibit the absorption of Enrofloxacin. It is recommended to separate both dosages for at least two hours.
- Enrofloxacin administered with theophylline can increase your serum levels. In dogs, theophylline clearance was reduced by 43% when co-administered with Enrofloxacin at a dose of 5 mg / kg every 24 hours.
- Concomitant administration with other drugs that have hepatic metabolism may affect the pharmacokinetics of one or both drugs.

CONTRAINDICATIONS

- Enrofloxacin is contraindicated in animals with hypersensitivity to quinolones.
- Do not administer to small to medium breed dogs, younger than 1 year, nor to large breed dogs younger than 1 ½ year.
- Do not administer to cats younger than 8 weeks of age.
- The use of Enrofloxacin can cause the formation of crystals in urine, therefore it is recommended not to administer in dehydrated animals.
- It is not recommended for use in pregnancy and / or lactation.

WARNINGS

- Mantener fuera del alcance de los niños.
- Las Fluoroquinolonas, como Enrofloxacino, no deben utilizarse como primera línea de tratamiento, excepto que no exista alternativa terapéutica disponible. Cuando se empleen como segundo tratamiento, deberá ser sobre la base de estudios de susceptibilidad.

ADVERSE EFFECTS

- Arthropathies have been reported in immature animals, especially dogs. The risk of arthropathy increases when the dose is increased, however, it has also been reported when the recommended doses are used in dogs.
- Other side effects that have been described in very rare cases (less than 1 animal in 10,000 animals, including isolated reports) include vomiting, anorexia, increased liver enzymes, ataxia, seizures, depression, lethargy, and nervousness.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
Discard the remains of unused product in its original container. Dispose of this product with caution with household waste.

CONSERVATION

Keep in a cool and dry place at room temperature between 15 ° and 30 ° C.

CONDITION OF SALE

For Chile: Sales under prescription Veterinary Doctor withheld.
For Export: Sale under Veterinary Medical prescription.

PRESENTATION

Case with 10 tablets.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 602
El Salvador: VE2013094805
Rep. Dominicana: Reg. Nº 8786
Bolivia: Reg. SENASAG PUV-F N° 007250/16
Costa Rica: Reg. N° MAG CL4-5-60-5759
Perú: Registro SENASA F.82.21.I.0385

COUNTRIES WHERE IT IS MARKETED

Distribution in El Salvador:
Rafael Alfredo Alfaro Castillo.
8a C. Pte. Pje. Moreno N ° 112, Col. Flor Blanca.
San Salvador, El Salvador.

Imported and distributed in Bolivia by:
ZOOFARMA

TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

Imported and Distributed in Peru by:
Representaciones Durand SAC.
Av. Manuel Olguín N ° 501 Office N ° 604 Santiago de Surco Lima.

ROSTRUM® 5% - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANTIBACTERIANO DE AMPLIO ESPECTRO.

Technical Specification

SPECIES

Bovines and pigs.

Rostrum® 5%, is a broad spectrum antibacterial, indicated for the treatment of bacterial infections caused by microorganisms sensitive to Enrofloxacin.
In cattle and pigs, its use is indicated in the treatment of respiratory infections, digestive system infections, Agalactia Mastitis Metritis Syndrome in sows, and septic conditions caused by the following bacterial species: *E. coli, Salmonella spp., Klebsiella spp. , Proteus, spp., Yersinia spp., Haemophilus spp., Pasteurella spp., Actinobacillus spp., Brucella spp., Campylobacter spp., Streptococcus spp, Staphylococcus spp., Bacteriodes spp., Bordetella spp. and Mycoplasma spp.*

DOSAGE FORM

Injectable solution

THERAPEUTIC ACTION

Broad spectrum antibacterial

COMPOSITION

Each 100 mL contains:
Enrofloxacin ..... 5 g
Excipients q.s. ..... 100 mL

INDICATIONS

- Do not administer Enrofloxacin to animals with hypersensitivity to Quinolones.
- The use of Enrofloxacin can cause the formation of crystals in urine, for which it is recommended not to administer in dehydrated animals.
- Do not use during pregnancy and lactation.
- Do not use in breeding animals.

ROUTE OD ADMINISTRATION AND DOSAGE

Route of administration:
Intramuscular (cattle and pigs) and subcutaneous (cattle).

Dose of the active principle:
In both species, 2.5 mg / Kg of weight, every 24 hours for 3 to 5 days. In acute infections, the dose can be increased to 5 mg / kg of weight, every 24 hours for 3 to 5 days.

Product dosage:
In both species, 1mL / 20 Kg of weight, every 24 hours for 3 to 5 days. In acute infections, the dose can be increased to 2 mL / 20 kg of weight, every 24 hours for 3 to 5 days.

DRUG INTERACTIONS

- Enrofloxacin administered with theophylline can increase your serum levels.
- Concomitant administration with other drugs that have hepatic metabolism may affect the pharmacokinetics of one or both drugs.

CONTRAINDICATIONS

- Do not administer Enrofloxacin to animals with hypersensitivity to Quinolones.
- The use of Enrofloxacin can cause the formation of crystals in urine, for which it is recommended not to administer in dehydrated animals.
- Do not use during pregnancy and lactation.
- Do not use in breeding animals.

PRECAUTIONS

- Avoid the use of Enrofloxacin in young growing animals due to the cartilaginous abnormalities that it can generate.
- Fluoroquinolones, like Enrofloxacin, should not be used as the first line of treatment, unless there is no therapeutic alternative available. When used as a second treatment, it should be based on susceptibility studies.

SPECIAL PRECAUTIONS FOR USE

- Avoid the use of Enrofloxacin in young growing animals due to the cartilaginous abnormalities that it can generate.
- Fluoroquinolones, like Enrofloxacin, should not be used as the first line of treatment, unless there is no therapeutic alternative available. When used as a second treatment, it should be based on susceptibility studies.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not handle by people hypersensitive to quinolones.
- In case of contact with skin, eyes or mucous membranes, wash immediately with plenty of water.
- In case of accidental injection, go immediately to a medical center and show the product insert

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

- Arthropathies have been reported in immature animals such as calves and piglets. The risk of arthropathy increases when the dose is increased.
- Other adverse effects that have been reported in very rare cases (less than 1 animal in 10,000 animals, including isolated reports) include vomiting, anorexia, increased liver enzymes, ataxia, seizures, depression, lethargy and nervousness.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be discarded as household waste, without any special precaution. Do not dispose of containers with product remains on the ground or watercourses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Storage conditions:
Store between 15 and 30 °C, protected from light. Once the container is opened, use within 4 weeks. Discard unused product after that period of time.

CONDITION OF SALE

Sale under retained Veterinary Medical prescription.

PRESENTATION

20 mL vial

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Reg. SAG N° 723

ROSTRUM® LARGE BREED - ORAL TABLET

COMPRIMIDO ORAL.

ANTIBACTERIANO.

Technical Specification

SPECIES

Dogs.

Rostrum® Large Breed, is a broad spectrum antibacterial, recommended for the treatment of:

- Gastrointestinal tract infections (*E. coli, Salmonella spp., Proteus spp.*).
- Respiratory tract infections (*Pasteurella spp., Bordetella spp., Klebsiella spp.*).
- Genital urinary infections such as nephritis, pyelonephritis, cystitis. (*E. coli, Corynebacterium pyogenes, Staphylococcus aureus, Streptococcus spp.*).
- Skin infections.
- Infections of the external auditory canal and otitis (*E. coli, Staphylococcus aureus, Streptococcus spp.*).
- Dermatitis, infected wounds (*E. coli, Klebsiella spp., Staphylococcus aureus, Streptococcus spp.*).

DOSAGE FORM

Oral Tablets.

THERAPEUTIC ACTION

Antibacterial.

COMPOSITION

Each tablet contains:
Enrofloxacin ..... 150 mg
Excipients q.s.p ..... 1 tablet

INDICATIONS

- Enrofloxacin is contraindicated in animals with hypersensitivity to quinolones.
- Do not administer to small to medium breed dogs, less than 1 year old, or to large breed dogs less than 1 ½ years old.
- Do not administer to animals with cartilaginous growth disorders.
- The use of Enrofloxacin can generate the formation of crystals in urine, for which it is recommended not to administer in dehydrated animals.
- Its use in pregnancy and / or lactation is not recommended.

ROUTE OD ADMINISTRATION AND DOSAGE

Orally.

- **Dose of the active principle:** 150 mg of Enrofloxacin / 30 Kg of weight per day for 7 to 10 days. Half the dose can be administered, twice a day.
- **Product dose:** 1 tablet of 150 mg / 30 Kg of weight per day for 7 to 10 days.

DRUG INTERACTIONS

- Compounds containing di or trivalent cations such as aluminum, calcium, magnesium, iron or zinc can decrease the absorption of Enrofloxacin. Thus, for example antacids containing cations (Mg2 +, Al3 +, Ca2 +) can bind to Enrofloxacin and prevent its absorption.
- Sucralfate can inhibit the absorption of Enrofloxacin. It is recommended to separate both dosages for at least two hours.
- Enrofloxacin administered with theophylline can increase your serum levels. In dogs, theophylline clearance was reduced by 43% when co-administered with Enrofloxacin at a dose of 5 mg / kg every 24 hours.
- Concomitant administration with other drugs that have hepatic metabolism may affect the pharmacokinetics of one or both drugs.

CONTRAINDICATIONS

- Enrofloxacin is contraindicated in animals with hypersensitivity to quinolones.
- Do not administer to small to medium breed dogs, less than 1 year old, or to large breed dogs less than 1 ½ years old.
- Do not administer to animals with cartilaginous growth disorders.
- The use of Enrofloxacin can generate the formation of crystals in urine, for which it is recommended not to administer in dehydrated animals.
- Its use in pregnancy and / or lactation is not recommended.

WARNINGS

- Mantener fuera del alcance de los niños.
- Las Fluoroquinolonas, como Enrofloxacino, no deben utilizarse como primera línea de tratamiento, excepto que no exista alternativa terapéutica disponible. Cuando se empleen como segundo tratamiento, deberá ser sobre la base de estudios de susceptibilidad.

ADVERSE EFFECTS

- Arthropathies have been reported in immature animals, especially dogs. The risk of arthropathy increases when the dose is increased, however, it has also been reported when the recommended doses are used in dogs.
- Other adverse effects that have been reported in very rare cases (less than 1 animal in 10,000 animals, including isolated reports) include vomiting, anorexia, increased liver enzymes, ataxia, seizures, depression, lethargy and nervousness.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Dispose of this product waste carefully with household waste.

CONSERVATION

Keep in a cool and dry place at room temperature between 15 and 30 ° C.

CONDITION OF SALE

For Chile: Sale with retained Veterinary Medical prescription.
For Export: Sale with Veterinary Medical prescription.

PRESENTATION

Box with 10 tablets.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N 721
Costa Rica: Reg. N° MAG CL-4-5-61-5699
El Salvador: VE2014034902
Bolivia: Reg. SENASAG PUV-F N° 007251/16

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

ROSTRUM® PLUS - OTIC EMULSION

EMULSIÓN ÓTICA, ANTIBACTERIANA Y ANTIMICÓTICA

ANTIBACTERIANO Y ANTIMICÓTICO.

Technical Specification

SPECIES

Dogs.

Indicated for the treatment of acute or chronic external otitis in dogs, caused by bacteria and fungi sensitive to Enrofloxacin and / or Silver Sulfadiazine.

DOSAGE FORM

Otic, antibacterial and antifungal emulsion

THERAPEUTIC ACTION

Antibacterial and antifungal

COMPOSITION

Each mL of emulsion contains:
Enrofloxacin ..... 5 mg
Silver Sulfadiazine ..... 10 mg
Excipients q.s.p ..... 1 mL

INDICATIONS

- Do not use in dogs with hypersensitivity to Quinolones and / or Sulfonamides.
- Do not use in dogs with perforated tympanic membrane.
- Do not use during pregnancy and lactation.

MODE OF APPLICATION

Clean the ear canal, then administer Rostrum® Plus in sufficient quantity to cover the external ear canal, gently massaging the ear to ensure a complete and even distribution of the medicine throughout the canal.

ROUTE OD ADMINISTRATION AND DOSAGE

Topical administration.

Product dosage: 5 to 10 drops in dogs weighing 15 kilos or less, and 10 to 15 drops in dogs weighing more than 15 kilos. Administer 2 times a day for 14 days.

DRUG INTERACTIONS

It is not recommended to use concomitantly with other pharmaceutical products.

CONTRAINDICATIONS

- Do not use in dogs with hypersensitivity to Quinolones and / or Sulfonamides.
- Do not use in dogs with perforated tympanic membrane.
- Do not use during pregnancy and lactation.

PRECAUTIONS

- The use of this combination has not been evaluated in dogs with perforation of the tympanic membrane. Consequently, before administering the product, the external auditory canal should be examined in order to ensure that there is no rupture of the tympanic membrane, in order to avoid contamination of the middle ear, as well as vestibular and cochlear lesions. If during the administration of the product there are signs of deafness or vestibular alterations, discontinue its use.
- Quinolones should be used with caution in animals with central nervous system (CNS) disorders. In these animals, in rare cases quinolones have been associated with CNS stimulation, which can generate seizures.
- Fluoroquinolones, such as Enrofloxacin, should not be used as the first line of treatment, unless there is no therapeutic alternative available. When used as a second treatment, it should be based on susceptibility studies.

SPECIAL PRECAUTIONS FOR USE

- The use of this combination has not been evaluated in dogs with perforation of the tympanic membrane. Consequently, before administering the product, the external auditory canal should be examined in order to ensure that there is no rupture of the tympanic membrane, in order to avoid contamination of the middle ear, as well as vestibular and cochlear lesions. If during the administration of the product there are signs of deafness or vestibular alterations, discontinue its use.
- Quinolones should be used with caution in animals with central nervous system (CNS) disorders. In these animals, in rare cases quinolones have been associated with CNS stimulation, which can generate seizures.
- Fluoroquinolones, such as Enrofloxacin, should not be used as the first line of treatment, unless there is no therapeutic alternative available. When used as a second treatment, it should be based on susceptibility studies.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Wash your hands after administering the product.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

- With the exception of possible cartilage abnormalities in young animals, the adverse effect profile of Enrofloxacin is generally limited to gastrointestinal problems (vomiting, anorexia), although rare incidents of elevated liver enzymes, ataxia, seizures, depression have also been reported. and lethargy.
- No evidence of systemic toxicity due to Silver Sulfadiazine has been reported in dogs, even 1% Silver Sulfadiazine ear suspensions have been used in dogs for more than three months without incident.

OBSERVATIONS

Shake before using.

Special precautions for disposal of unused product or waste material:
Dispose of any unused product remains in its original, tightly closed container. Dispose of this product waste carefully with household waste.

CONSERVATION

Keep in a cool place, at a temperature between 15° and 30° C, protected from light.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

Dropper bottle with 15 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 1523
Bolivia: Reg. SENASAG PUV-F-N° 006147/14
El Salvador: VE2014084963

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:
ZOOFARMA

TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

Distributed in El Salvador by:
Rafael Alfredo Alfaro Castillo.

Imported and Distributed in Peru by:
Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

RUMITEN® ORAL SUSPENSION

SUSPENSIÓN ORAL.

ANTIPARASITARIO DE AMPLIO ESPECTRO.

Technical Specification

SPECIES

Sheep and goats.

Indicated for the treatment of gastrointestinal and pulmonary parasitosis in sheep and goats caused by parasites sensitive to the combination of Fenbendazole and Praziquantel, such as: *Moniezia spp.*, Mature and immature forms of *Haemochus spp., Cooperia spp., Nematodirus spp., Chabertia ovina , Oesophagostomum spp., Trichostrogylus spp., Trichuris ovis., Ostertagia ostertagi* (hypobiotic adults and larvae), *Ostertagia circumcincta* and *Dictyocaulus filaria*. It is recommended to treat all the animals in the herd every three months. This contributes to reducing the pressure of parasitic infection, achieving epidemiological control.

DOSAGE FORM

Oral suspension

THERAPEUTIC ACTION

Broad spectrum antiparasitic

COMPOSITION

Each 100 mL of oral suspension contains:
Fenbendazole ..... 2 g
Praziquantel ..... 1.5 g
Excipients q.s.p ..... 100 mL

INDICATIONS

Do not administer to animals with known hypersensitivity to any active ingredient.

EFFECTIVENESS

Efficacy period: 36 months.

ROUTE OD ADMINISTRATION AND DOSAGE

Oral administration.

Recommended practice dose:
2.5 mL per 10 kg of body weight in a single dose (equivalent to a dose of 3.75 mg of Praziquantel and 5 mg of Fenbendazole per kg of live weight).

CONTRAINDICATIONS

Do not administer to animals with known hypersensitivity to any active ingredient.

WARNINGS

- Agitar antes de usar.
- Mantener alejado del alcance de los niños.
- Solamente para uso oral en ovinos y caprinos.

GUARD PERIOD

Meat: 14 days.

Milk: 7 days.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of natural water. Do not reuse the container. Dispose of this product waste carefully with household waste.

CONSERVATION

Store at a temperature between 2 and 30 ° C, protected from light.
Once the container is opened, use within 1 month. Discard the unused product after that period.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Bottle with 100 mL, 1 Liter and 3 Liters.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Reg. SAG Nº 1961

SENILPET® CEREBRAL 20 ORAL TABLET.

COMPRIMIDO ORAL.

SUPLEMENTO NUTRICIONAL PARA PERROS DE EDAD AVANZADA

Technical Specification

SPECIES

Senior dogs (+ 7 years)
Large breeds

SENILPET® CEREBRAL 20 is a nutritional supplement for older dogs and large breeds, which has been specially formulated to support brain function and cognitive health of your pet.

SENILPET® CEREBRAL 20 provides nutritional support to your pet through compounds that help protect it from normal aging, improving your dog's activity levels and vitality.

DOSAGE FORM

Oral tablet.

THERAPEUTIC ACTION

Nutritional supplement for elderly dogs.

COMPOSITION

Each tablet contains:
Phosphatidylserine ..... 50 mg
Ginkgo biloba ..... 100 mg
Vitamin B6 (Pyridoxine HCl) ..... 41.2 mg
Vitamin E (D-alpha Tocopherol) ..... 67 IU
Excipients c.s.p ..... 1 tablet

INGREDIENTS

Palatable agent based on chicken giblets, Vitamin E, Phosphatidylserine, Ginkgo biloba, Vitamin B6 and authorized preservatives.

PROPERTIES

Supports your pet's brain function and cognitive health

- For older dogs (+7)
- Helps improve activity levels and vitality of pets
- Highly palatable tablets
- For Large breed dogs

Senilpet® Cerebral 5 contains:

- **Phosphatidylserine:** Membrane phospholipid that helps improve communication between nerve cells, through the synthesis and release of neurotransmitters.
- **Ginkgo biloba:** Helps improve cerebral blood flow, increases glucose metabolism, and has a powerful antioxidant effect.
- **Pyridoxine (Vitamin B6):** It is essential to help maintain brain health and function. Pyridoxine is a cofactor of the biosynthesis of neurotransmitters such as Serotonin, Noradrenaline and Dopamine.
- **Vitamin E (D-alpha Tocopherol):** Powerful antioxidant that provides protection against oxidative stress naturally generated in the cellular aging process.

INDICATIONS

Contraindicated in animals receiving anticoagulant drugs.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer 1 tablet of SENILPET® CEREBRAL 20 for every 20 kilos of body weight, once a day.
SENILPET® CEREBRAL 20 has been formulated as a nutritional supplement for use for long periods of time. For best results, give the recommended dose for at least 1 to 2 months.

CONTRAINDICATIONS

Contraindicated in animals receiving anticoagulant drugs.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

VETERINARY USE.
EXCLUSIVE USE IN ANIMAL FEED.
IT DOES NOT CORRESPOND TO A COMPLETE FOOD.

CONSERVATION

Keep in a cool, dry place and protected from light, at no more than 30 ° C.

CONDITION OF SALE

OTC product

PRESENTATION

Bottle with 30 tablets

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. LENAA N°: RM 03-008N
Perú: Reg. SENASA A.36.15.I.0573
Costa Rica: Lic. DAA-MAG 579-026

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Peru by Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

SENILPET® CEREBRAL 5 ORAL TABLET.

COMPRIMIDO ORAL.

SUPLEMENTO NUTRICIONAL PARA PERROS DE EDAD AVANZADA.

Technical Specification

SPECIES

Senior dogs (+ 7 years)
Small breeds

SENILPET® CEREBRAL 5, is a nutritional supplement for elderly dogs, which has been specially formulated to support your pet's brain function and cognitive health.

SENILPET® CEREBRAL 5 provides nutritional support to your pet through compounds that help protect it from normal aging, improving the levels of activity and vitality of your dog.

DOSAGE FORM

Oral tablet.

THERAPEUTIC ACTION

Nutritional supplement for elderly dogs.

COMPOSITION

Each tablet contains:
Phosphatidylserine ..... 12.5 mg
Ginkgo biloba ..... 25.0 mg
Vitamin B6 (Pyridoxine HCL) ..... 10.3 mg
Vitamin E (D-alpha Tocopherol) ..... 8.4 IU
Excipients q.s.p ..... 1 tablet

INGREDIENTS

Palatable agent based on chicken giblets, Vitamin E, Phosphatidylserine, Ginkgo biloba, Vitamin B6 and authorized preservatives.

PROPERTIES

Supports your pet's brain function and cognitive health

- For older dogs (+7)
- Helps improve activity levels and vitality of pets
- Highly palatable tablets

Senilpet® Cerebral 5 contains:

- **Phosphatidylserine:** Membrane phospholipid that helps improve communication between nerve cells, through the synthesis and release of neurotransmitters.
- **Ginkgo biloba:** Helps improve cerebral blood flow, increases glucose metabolism, and has a powerful antioxidant effect.
- **Pyridoxine (Vitamin B6):** It is essential to help maintain brain health and function. Pyridoxine is a cofactor of the biosynthesis of neurotransmitters such as Serotonin, Noradrenaline and Dopamine.
- **Vitamin E (D-alpha Tocopherol):** Powerful antioxidant that provides protection against oxidative stress naturally generated in the cellular aging process.

INDICATIONS

Contraindicated in animals receiving anticoagulant drugs.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer 1 tablet of SENILPET® CEREBRAL 5 every 5 kilos of body weight, once a day.
SENILPET® CEREBRAL 5 has been formulated as a nutritional supplement for use for long periods of time. For best results, give the recommended dose for at least 1 to 2 months.

CONTRAINDICATIONS

Contraindicated in animals receiving anticoagulant drugs.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

VETERINARY USE.
EXCLUSIVE USE IN ANIMAL FEED.
IT DOES NOT CORRESPOND TO A COMPLETE FOOD.

CONSERVATION

Keep in a cool, dry place and protected from light, at no more than 30 ° C.

CONDITION OF SALE

OTC product

PRESENTATION

Bottle with 60 tablets

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. LENAA N°: RM 03-008N
Perú: Reg. SENASA A.36.15.I.0574
Costa Rica: Reg. Lic. DAA-MAG 579-027

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Peru by Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

SILIMADRAG® ORAL SUSPENSION.

SUSPENSIÓN ORAL.

SUPLEMENTO NUTRICIONAL PARA PERROS Y GATOS.

Technical Specification

SPECIES

Dogs and cats

Silimadrag® is a nutritional supplement of natural origin, specially formulated to maintain the liver health of your pet and support states that occur with liver function disorders of various origins (medicinal, infectious or toxic, among others).

DOSAGE FORM

Oral suspension.

THERAPEUTIC ACTION

Nutritional supplement for dogs and cats.

COMPOSITION

Each mL of suspension contains:
Silymarin ....25 mg
DL-Methionine ..... .80 mg
Choline Chloride .... 80 mg
Excipients q.s.p .... 1 mL

PROPERTIES

Silymarin is a flavonolignan, extracted from the fruit of milk thistle, which has various cytoprotective and regenerative properties that modulate liver function and promote a hepatoprotective effect. Methionine is an essential amino acid that participates in metabolism and in hepatic cytoprotective mechanisms. Choline, for its part, is an essential vitamin that plays an important role in fat metabolism, being a donor of functional groups that participate in the antioxidant system of the liver.

ROUTE OD ADMINISTRATION AND DOSAGE

Orally

- Dogs: 4 mL every 10 kg of weight, 2 times a day.
- Cats: 1 mL every 2.5 kg of weight, 2 times a day.

For optimal results, give the suggested dose for at least 1 to 2 months.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

- exclusive use in animal feed
- does not correspond to a complete food

CONSERVATION

Keep in a cool, dry place and away from light, at no more than 30°C.

CONDITION OF SALE

Free sale without prescription

PRESENTATION

Bottle with 120 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Reg. LENAA N°: RM 03-008N

Costa Rica: Reg. Lic. DAA-MAG 579-028

SINPUL® SHAMPOO

SHAMPOO.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Dogs.

External antiparasitic shampoo for action against fleas (*Ctenocephalides felis, Ctenocephalides canis*) and ticks (*Rhipicephalus sanguineus*) in dogs.

DOSAGE FORM

Shampoo.

THERAPEUTIC ACTION

External Antiparasitics.

COMPOSITION

Each 100 mL of shampoo contains:
Permethrin.....1.0 g
Propoxur.....0.1 g
Excipients q.s.....100 mL

INDICATIONS

- Do not administer to puppies younger than 3 months.
- Do not administer to animals hypersensitive to pyrethroids or carbamates.
- Do not administer to pregnant or lactating females.

MODE OF APPLICATION

- Wet your pet's coat with warm water and apply enough shampoo on the coat.
- Massage until abundant foam is obtained; leave to act for 3 to 5 minutes and then rinse.
- Repeat the operation and then dry the pet.

ROUTE OD ADMINISTRATION AND DOSAGE

For topical use.

CONTRAINDICATIONS

- Do not administer to puppies younger than 3 months.
- Do not administer to animals hypersensitive to pyrethroids or carbamates.
- Do not administer to pregnant or lactating females.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Wear gloves when applying the product.
- Wash your hands with soap and water after handling the product.
- In the event that there is contact with the skin and / or the eyes, it is recommended to immediately wash with plenty of water.
- In case of accidental ingestion it is recommended to rinse

ANTIDOTE

Atropine sulfate.

WARNINGS

Advertencias y precauciones especiales de uso:

- Mantener fuera del alcance de los niños.
- En caso de ingestión accidental por el animal consulte a su Médico Veterinario.
- Antídoto: Atropina.

OBSERVATIONS

Special precautions for the disposal of unused product or waste material:
Empty containers can be discarded as household waste, without any special precautions. Do not dispose of containers with product residues on the ground or water courses. For expired or unused products contact the manufacturer laboratory.

CONSERVATION

Store between 15 and 30 ° C, protected from light.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

Bottle with 300 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No. 934

SINPUL® SPRAY SOLUTION

SOLUCIÓN SPRAY.

ANTIPARASITARIO EXTERNO

Technical Specification

SPECIES

Dogs.

Eliminates fleas (*Ctenocephalides felis*) and ticks (*Rhipicephalus sanguineus*) in dogs.

DOSAGE FORM

Spray solution.

THERAPEUTIC ACTION

External Antiparasitics.

COMPOSITION

Each 100 mL contains:
Permethrin.....0.2 g
Piperonyl Butoxide.....2.0 g
Excipients q.s.....100.0 mL

INDICATIONS

- Do not use in cats.
- Do not use in puppies under 6 weeks of age.
- Do not administer in animals with hypersensitivity to the active principles.
- Do not administer in pregnant or lactating females.

MODE OF APPLICATION

Spray the product all over the dog's body, from the tail to the neck (application against the grain). After applying, rub the entire body vigorously with your hands so that the product achieves maximum contact with the skin and not only remains on the coat. If the dog has a large number of fleas and / or ticks, it is recommended to repeat the application every 7 to 15 days.

ROUTE OD ADMINISTRATION AND DOSAGE

Topical use.

Dosage of active ingredients:
6 to 18 mg / Kg of Permethrin and 60 to 80 mg / Kg of Piperonyl Butoxide.

Product dosage:
Apply 4 to 11 sprays per kg of weight, depending on the length of the coat, in a single dose.

CONTRAINDICATIONS

- Do not use in cats.
- Do not use in puppies under 6 weeks of age.
- Do not administer in animals with hypersensitivity to the active principles.
- Do not administer in pregnant or lactating females.

PRECAUTIONS

- Flammable product.
- Avoid contact with the eyes, mouth or other mucous membranes.
- In case of irritation, stop using the product and wash with plenty of water.

SPECIAL PRECAUTIONS FOR USE

- Flammable product.
- Avoid contact with the eyes, mouth or other mucous membranes.
- In case of irritation, stop using the product and wash with plenty of water.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- When applying the product wear protective gloves.
- Wash your hands with plenty of water after use.
- In case of contact with the eyes, mouth or skin, wash the affected area with plenty of water.

WARNINGS

- Mantener fuera del alcance de los niños.
- Uso externo.

ADVERSE EFFECTS

Hypersensitivity reactions have been described when Permethrin is administered topically. In some cases, itching or mild irritation may occur at the application site.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard the remains of unused product in its original container, well closed. Do not throw the empty container or with product remains, in soils, rivers, lakes or streams of water, as it is toxic to aquatic animals. Do not reuse the container. Dispose of this product waste carefully with household waste.

CONSERVATION

Keep at room temperature (between 15 and 30 ° C), in a dry place and away from fire. Once opened, use the product within 12 weeks. Discard the unused product after that period of time.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

200 mL and 500 mL spray bottle

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG N° 0390

SINPULDRY® CAT - TOPICAL POWDER

POLVO TÓPICO.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Cats.

Eliminate fleas and ticks on cats.

DOSAGE FORM

Topical powder.

THERAPEUTIC ACTION

External Antiparasitics.

COMPOSITION

Each 100 g contains:

Carbaryl.....2 g

Excipients q.s..... 100 g

INDICATIONS

- Do not administer to cats under 6 weeks of age
- Do not administer in pregnant or lactating females
- Do not use in animals with known hypersensitivity to Carbaryl

MODE OF APPLICATION

Sprinkle enough of the product on the fur, preferably in the area of the neck and back and rubbing to get to the skin. For a better application, brush the fur to spread the powder over the entire body surface.

ROUTE OD ADMINISTRATION AND DOSAGE

For topical use.

CONTRAINDICATIONS

- Do not administer to cats under 6 weeks of age
- Do not administer in pregnant or lactating females
- Do not use in animals with known hypersensitivity to Carbaryl

PRECAUTIONS

- Remove excess product from the animal
- Avoid contact of the product with eyes, mouth and mucous membranes of the animal
- Avoid the animal ingesting the product
- Animals with low cholinesterase activity are more sensitive to inhibition of it by carbamates derivatives
- Long hair thin cats are more susceptible to cholinesterase inhibition due to lack of fat

SPECIAL PRECAUTIONS FOR USE

- Remove excess product from the animal
- Avoid contact of the product with eyes, mouth and mucous membranes of the animal
- Avoid the animal ingesting the product
- Animals with low cholinesterase activity are more sensitive to inhibition of it by carbamates derivatives
- Long hair thin cats are more susceptible to cholinesterase inhibition due to lack of fat

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Avoid contact of the product with eyes, mouth or other mucous membranes
- In case of irritation, wash with plenty of water
- The use of gloves and a mask is recommended during the application
- Wash hands thoroughly after application of the product
- Keep the container closed after use

WARNINGS

- En caso de ingesta accidental, consulte al médico.
- Mantener el envase cerrado y fuera del alcance de los niños
- Uso externo.
- VENENO.

CONSERVATION

Keep in a fresh, dry place, at room temperature, between 15 and 30 °C.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

100 g

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No.: 0414

SINPULDRY® DOG - TOPICAL POWDER

POLVO TÓPICO.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Dogs.

Eliminate fleas and ticks on dogs.

DOSAGE FORM

Topical powder.

THERAPEUTIC ACTION

External antiparasitic.

COMPOSITION

Each 100 g contains:

Carbaryl.....2 g

Excipients q.s.....100 g

INDICATIONS

- Do not administer to dogs under 4 weeks of age
- Do not use in pregnant or lactating animals
- Do not use in animals with known hypersensitivity to Carbaryl

MODE OF APPLICATION

Sprinkle over the fur, preferably on the kneck and flank, by rubbing until the skin has been reached.

ROUTE OD ADMINISTRATION AND DOSAGE

For topical use.

CONTRAINDICATIONS

- Do not administer to dogs under 4 weeks of age
- Do not use in pregnant or lactating animals
- Do not use in animals with known hypersensitivity to Carbaryl

PRECAUTIONS

SPECIAL PRECAUTIONS FOR USE

SPECIAL PRECAUTIONS FOR THE OPERATOR

Precautions for the product handler:

- Avoid contact of the product with eyes, mouth or other mucous membranes
- In case of irritation, wash with plenty of water
- The use of gloves and a mask is recommended during the application
- Wash hands thoroughly after application of the product
- Keep the container closed after use

WARNINGS

- Uso externo.
- En caso de ingesta accidental, consulte al médico.
- Mantener el envase cerrado y fuera del alcance de los niños.
- VENENO.

CONSERVATION

Keep in a fresh, dry place, at room temperature, between 15 and 30 °C.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

100 g

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Reg. SAG No.: 0414

SINPULKILL® CATS - COLLAR

COLLAR.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Cats.

Antiparasitic collar effective in the treatment and control of flea infestation (*Ctenocephalides felis*) in cats, guaranteeing flea control for 4 months.

DOSAGE FORM

Collar.

THERAPEUTIC ACTION

External Antiparasitic.

COMPOSITION

Each 100 g of collar contains:
Diazinon .... 15.0 g
Excipients q.s.p ..... 100.0 g

INDICATIONS

- Do not use in sick or convalescent animals.
- Do not use in cats under 6 months of age.
- Do not use in Persian cats.
- Do not administer to animals hypersensitive to organophosphate compounds, such as Diazinon.
- Do not use during pregnancy, lactation or in breeding animals.

USE INSTRUCTIONS

Attach the collar loosely to the cat's neck and cut off the remaining collar that protrudes from the buckle. It is recommended to leave a space of two fingers between the collar and the neck.

DRUG INTERACTIONS

- It is recommended not to use organophosphates together with other cholinesterase inhibitor compounds, especially antiparasitics that use such a mechanism of action.
- Do not use concomitantly with other antiparasitic products.

CONTRAINDICATIONS

- Do not use in sick or convalescent animals.
- Do not use in cats under 6 months of age.
- Do not use in Persian cats.
- Do not administer to animals hypersensitive to organophosphate compounds, such as Diazinon.
- Do not use during pregnancy, lactation or in breeding animals.

PRECAUTIONS

- Prevent the animal from playing, licking or swallowing the collar.
- Remove the collar in case of irritation in the contact area, or if signs such as salivation, vomiting or diarrhea appear.
- For external use only.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR USE

- Prevent the animal from playing, licking or swallowing the collar.
- Remove the collar in case of irritation in the contact area, or if signs such as salivation, vomiting or diarrhea appear.
- For external use only.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not handle by people who are hypersensitive to organophosphate compounds, such as Diazinon.
- Do not handle by pregnant or lactating women.
- Wear gloves when handling the collar, either when putting it on or taking it off.
- Do not smoke, eat or drink near the collar.
- In case of accidental ingestion, go immediately to a medical center and show the product label.
- Prevent the owner or children from sleeping with the animal while it is wearing the collar.
- Wash your hands after handling the collar.
- Prevent children from playing with the collar.

ADVERSE EFFECTS

In very small cases there may be irritation in the contact area.

OBSERVATIONS

It not contains metallic parts for preventing allergies.

Special precautions for disposal of unused product or waste material:

- Empty containers can be disposed of as household waste, without any special precautions.
- Do not dispose of containers with product remains on the ground or water courses.
- For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 ° C, protected from light.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

Case with 1 collar.

PREPARED BY

Manufactured by:Societé AB7 Industries S.A. Chemin des Monges Deyme, 314150 Montgiscard. France.
Imported and distributed by:
Laboratorio Drag Pharma Chile Invetec S.A. Lautaro N ° 300, Quilicura. Santiago. Chile, under license from Societé AB7 Industries S.A. France.

RECORDS

Reg. S.A.G. No.: 1647

SINPULKILL® DOG COLLAR

COLLAR.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Dogs

Effective antiparasitic collar in the treatment and control of infestation by fleas (*Ctenocephalides felis*) and ticks (*Rhipicephalus sanguineus*) in dogs, guaranteeing flea control for 4 months, and ticks for 3 months.

DOSAGE FORM

Collar.

THERAPEUTIC ACTION

External Antiparasitics.

COMPOSITION

Each 100 g of collar contains:
Diazinon .... 15.0 g
Excipients q.s.p ..... 100.0 g

INDICATIONS

- Do not use in sick or convalescent animals.
- Do not use in pregnant, lactating or breeding animals.
- Do not use in dogs under 6 months of age.
- Do not administer to animals hypersensitive to organophosphate compounds, such as Diazinon, preferably Greyhound and Whippet breeds.

USE INSTRUCTIONS

Attach the collar loosely to the dog's neck and cut off the remaining collar protruding from the buckle. It is recommended to leave a space of two fingers between the collar and the neck.

DRUG INTERACTIONS

- It is recommended not to use organophosphates together with other cholinesterase inhibitor compounds, especially antiparasitics that use such a mechanism of action.
- Do not use concomitantly with other antiparasitic products.

CONTRAINDICATIONS

- Do not use in sick or convalescent animals.
- Do not use in pregnant, lactating or breeding animals.
- Do not use in dogs under 6 months of age.
- Do not administer to animals hypersensitive to organophosphate compounds, such as Diazinon, preferably Greyhound and Whippet breeds.

PRECAUTIONS

- Prevent the animal from playing, licking or swallowing the collar.
- Remove the collar in case of irritation in the contact area, or if signs such as salivation, vomiting or diarrhea appear.
- For external use only.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR USE

- Prevent the animal from playing, licking or swallowing the collar.
- Remove the collar in case of irritation in the contact area, or if signs such as salivation, vomiting or diarrhea appear.
- For external use only.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not handle by people who are hypersensitive to organophosphate compounds, such as Diazinon.
- Do not handle by pregnant or lactating women.
- Wear gloves when handling the collar, either when putting it on or taking it off.
- Do not smoke, eat or drink near the collar.
- In case of accidental ingestion, go immediately to a medical center and show the product label.
- Prevent the owner or children from sleeping with the animal while it is wearing the collar.
- Wash your hands after handling the collar.
- Prevent children from playing with the collar.

OBSERVATIONS

It not contains metallic parts for preventing allergies.

Special precautions for disposal of unused product or waste material:

- Empty containers can be disposed of as household waste, without any special precautions.
- Do not dispose of containers with product remains on the ground or water courses.
- For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 and 30 ° C, protected from light.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

Case with 1 Collar

PREPARED BY

Manufactured by: Societé AB7 Industries S.A. Chemin des Monges Deyme, 314150 Montgiscard. France.
Exported by:
Vetpharma Animal Health S.L. Les Corts, 23. 08028 Barcelona. Spain.
Imported and distributed in Chile by Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Reg. S.A.G. No.: 1646

SINPULSPOT® PLUS LARGE BREED - SPOT ON

SOLUCIÓN TÓPICA.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Dogs over 15 Kg of body weight.

Effective insecticide in the control and prevention of infestation by fleas (*Ctenocephalides felis*) and ticks (*Rhipicephalus sanguineus*) in dogs. The product provides effective protection against reinfestations for 4 weeks after application.

DOSAGE FORM

Topical solution. Spot on

THERAPEUTIC ACTION

External antiparasitic. Spot-on

COMPOSITION

Each dose of 2.0 mL contains:
Permethrin (cis/trans 25/75)..... 650 mg
Piperonyl Butoxide.....650 mg
Methoprene.....75 mg
Excipients q.s.....2.0 mL

INDICATIONS

- Do not apply to dogs younger than 4 months.
- **Do not apply to cats.**
- Do not apply to pregnant or lactating females.

MODE OF APPLICATION

Cut off the tip of the pipette (s). Separate the hair in the area of application, so that the dose is administered directly to the skin.
- In dogs over 15 kg apply a 2 mL pipette at the withers and another 2 mL pipette at the base of the tail (SINPULSPOT® PLUS, Large Breed).

ROUTE OD ADMINISTRATION AND DOSAGE

Dose: 4 mL (2 pipettes).
Single dose. Apply the recommended dose every 4 weeks.

DRUG INTERACTIONS

Do not use concomitantly with other pharmaceuticals.

CONTRAINDICATIONS

- Do not apply to dogs younger than 4 months.
- **Do not apply to cats.**
- Do not apply to pregnant or lactating females.

PRECAUTIONS

Special warnings and precautions for use:

- Keep out of the reach of children.
- For a greater residual effect, it is recommended to apply SINPULSPOT® PLUS, 3 days after the last bath of your dog, without re-bathing until it is strictly necessary.

SPECIAL PRECAUTIONS FOR USE

Special warnings and precautions for use:

- Keep out of the reach of children.
- For a greater residual effect, it is recommended to apply SINPULSPOT® PLUS, 3 days after the last bath of your dog, without re-bathing until it is strictly necessary.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Avoid contact with skin: wash your hands after handling the product.
- Avoid contact with eyes and in case of splashes wash with plenty of water.

ADVERSE EFFECTS

Unwanted effects and adverse reactions:
On very rare occasions, slight coloration may occur at the application site in light-coated dogs; transient signs of skin irritation such as redness, erythema, or itching; behavioral changes such as restlessness; and / or gastrointestinal signs such as salivation, vomiting or loss of appetite, which should disappear 24 hours after application. If signs of irritation persist, it is recommended to bathe the dog with a mild shampoo to remove the product and rinse with plenty of water. If the signs described continue or worsen, consult a Veterinary Doctor.

OBSERVATIONS

External use only.
Poison

Special precautions for the disposal of unused product or waste material:
Discard the remains of unused product in its original container. Do not discard the empty container or with product residues in rivers, lakes or streams of natural water. The product is toxic to aquatic organisms. Dispose of this product with caution with household waste.

CONSERVATION

Keep in a fresh, dry place, at room temperature between 2 and 30 °C.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

2 pipettes of 2 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG Nº: 1672
Costa Rica: Reg. N° MAG CL4-43-04-4139

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SINPULSPOT® PLUS MEDIUM BREED - SPOT ON

SOLUCIÓN TÓPICA.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Dog between 5 Kg and 15 Kg of body weight.

Effective insecticide in the control and prevention of infestation by fleas (*Ctenocephalides felis*) and ticks (*Rhipicephalus sanguineus*) in dogs. The product provides effective protection against reinfestations for 4 weeks after application.

DOSAGE FORM

Topical solution. Spot on

THERAPEUTIC ACTION

External antiparasitic. Spot on

COMPOSITION

Each dose of 2.0 mL contains:
Permethrin (cis/trans 25/75).....650 mg
Piperonyl Butoxide.....650 mg
Methoprene.....75 mg
Excipients q.s.....2.0 mL

INDICATIONS

- Do not apply to dogs younger than 4 months.
- **Do not apply to cats.**
- Do not apply to pregnant or lactating females.

MODE OF APPLICATION

Cut off the tip of the pipette (s). Separate the hair in the area of application, so that the dose is administered directly to the skin.

- In dogs from 5 to 15 kg apply the entire content of a 2 mL pipette (SINPULSPOT® PLUS, Medium Breed) at the withers (on the back between the shoulders).

ROUTE OD ADMINISTRATION AND DOSAGE

Dose: 2 mL
Single dose. Apply the recommended dose every 4 weeks.

DRUG INTERACTIONS

Do not use concomitantly with other pharmaceuticals.

CONTRAINDICATIONS

- Do not apply to dogs younger than 4 months.
- **Do not apply to cats.**
- Do not apply to pregnant or lactating females.

PRECAUTIONS

Special warnings and precautions for use:

- Keep out of the reach of children.
- For a greater residual effect, it is recommended to apply SINPULSPOT® PLUS, 3 days after the last bath of your dog, without re-bathing until it is strictly necessary.

SPECIAL PRECAUTIONS FOR USE

Special warnings and precautions for use:

- Keep out of the reach of children.
- For a greater residual effect, it is recommended to apply SINPULSPOT® PLUS, 3 days after the last bath of your dog, without re-bathing until it is strictly necessary.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Avoid contact with skin: wash your hands after handling the product.
- Avoid contact with eyes and in case of splashes wash with plenty of water.

ADVERSE EFFECTS

Unwanted effects and adverse reactions:
On very rare occasions, slight coloration may occur at the application site in light-coated dogs; transient signs of skin irritation such as redness, erythema, or itching; behavioral changes such as restlessness; and / or gastrointestinal signs such as salivation, vomiting or loss of appetite, which should disappear 24 hours after application. If signs of irritation persist, it is recommended to bathe the dog with a mild shampoo to remove the product and rinse with plenty of water. If the signs described continue or worsen, consult a Veterinary Doctor.

OBSERVATIONS

External use only.
Poison

Special precautions for the disposal of unused product or waste material:
Discard the remains of unused product in its original container. Do not discard the empty container or with product residues in rivers, lakes or streams of natural water. The product is toxic to aquatic organisms. Dispose of this product with caution with household waste.

CONSERVATION

Keep in a fresh, dry place, at room temperature between 2 and 30 °C.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

1 pipette of 2 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG Nº: 1672
Costa Rica: Reg. N° MAG CL4-43-04-4139

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SINPULSPOT® PLUS SMALL BREED - SPOT ON

SOLUCIÓN TÓPICA.

ANTIPARASITARIO EXTERNO.

Technical Specification

SPECIES

Dogs up to 5 Kg of body weight.

Effective insecticide in the control and prevention of infestation by fleas (*Ctenocephalides felis*) and ticks (*Rhipicephalus sanguineus*) in dogs. The product provides effective protection against reinfestations for 4 weeks after application.

DOSAGE FORM

Topic solution. Spot on

THERAPEUTIC ACTION

External antiparasitic. Spot on

COMPOSITION

Each dose of 1.0 mL contains:
Permethrin (cis/trans 25/75).....325 mg
Piperonyl Butoxide.....325 mg
Methoprene.....37.5 mg
Excipients q.s.....1.0 mL

INDICATIONS

- Do not apply to dogs younger than 4 months.
- **Do not apply to cats.**
- Do not apply to pregnant or lactating females.

MODE OF APPLICATION

Cut off the tip of the pipette (s). Separate the hair in the area of application, so that the dose is administered directly to the skin.

- In dogs weighing less than 5 kg, apply the entire content of a 1 mL pipette (SINPULSPOT® PLUS, Small Breed) at the withers (on the back between the shoulders).

ROUTE OD ADMINISTRATION AND DOSAGE

Dose: 1 mL
Single dose. Apply the recommended dose every 4 weeks.

DRUG INTERACTIONS

Do not use concomitantly with other pharmaceuticals.

CONTRAINDICATIONS

- Do not apply to dogs younger than 4 months.
- **Do not apply to cats.**
- Do not apply to pregnant or lactating females.

PRECAUTIONS

Special warnings and precautions for use:

- Keep out of the reach of children.
- For a greater residual effect, it is recommended to apply SINPULSPOT® PLUS, 3 days after the last bath of your dog, without re-bathing until it is strictly necessary.

SPECIAL PRECAUTIONS FOR USE

Special warnings and precautions for use:

- Keep out of the reach of children.
- For a greater residual effect, it is recommended to apply SINPULSPOT® PLUS, 3 days after the last bath of your dog, without re-bathing until it is strictly necessary.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Avoid contact with skin: wash your hands after handling the product.
- Avoid contact with eyes and in case of splashes wash with plenty of water.

ADVERSE EFFECTS

Unwanted effects and adverse reactions:
On very rare occasions, slight coloration may occur at the application site in light-coated dogs; transient signs of skin irritation such as redness, erythema, or itching; behavioral changes such as restlessness; and / or gastrointestinal signs such as salivation, vomiting or loss of appetite, which should disappear 24 hours after application. If signs of irritation persist, it is recommended to bathe the dog with a mild shampoo to remove the product and rinse with plenty of water. If the signs described continue or worsen, consult a Veterinary Doctor.

OBSERVATIONS

External use only.
Poison

Special precautions for the disposal of unused product or waste material:
Discard the remains of unused product in its original container. Do not discard the empty container or with product residues in rivers, lakes or streams of natural water. The product is toxic to aquatic organisms. Dispose of this product with caution with household waste.

CONSERVATION

Keep in a fresh, dry place, at room temperature between 2 and 30°C.

CONDITION OF SALE

OTC product (non-prescription)

PRESENTATION

1 pipette of 1 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. SAG Nº: 1672
Costa Rica: Reg. N° MAG CL4-43-04-4139

AVAILABLE FOR SALE

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SIR DOG® BLACK - SHAMPOO

SHAMPOO.

SHAMPOO PARA PERROS DE PELAJE NEGRO.

Technical Specification

SPECIES

Dogs.

SIR DOG® BLACK is specially formulated to enhance the black color of your pet's coat. Its composition, enriched with Aloe Vera Extract, favors a healthy and glossy coat.

DOSAGE FORM

Shampoo.

THERAPEUTIC ACTION

Black coat dog shampoo.

INGREDIENTS

Purified water, Lauryl ether sulfate sodium salt, Cocamidopropyl betaine, Cocamide DEA, Propylene glycol, Glycol distearate and laureth-4, Sodium Chloride, Aloe Vera extract, Formaldehyde 37%, Black colorant, Fragrance, EDTA disodium salt, Citric acid anhydrous.

PROPERTIES

Black coat dog shampoo with Aloe Vera extract

USE INSTRUCTIONS

- Wet your dog's coat completely with warm water.
- Apply SIR DOG® BLACK in a quantity enough to reach the skin.
- Massage until obtaining abundant lather, and let act for 3 to 5 minutes.
- Rinse with plenty of warm water.

SPECIAL PRECAUTIONS FOR THE OPERATOR

The use of gloves is recommended for preventing skin coloration of the person applying the product.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

For external use.

CONDITION OF SALE

OTC product.

PRESENTATION

390 mL

PREPARED BY

Drag Pharma Laboratory.

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SIR DOG® CONDITIONER

ACONDICIONADOR.

ACONDICIONADOR.

Technical Specification

SPECIES

Dogs.

SIR DOG® CONDITIONER is specially formulated to provide softness and elegance to your pet's coat. Unlike other conditioners, SIR DOG® CONDITIONER provides a maximum conditioning effect, without causing skin irritation or dryness. Its composition allows to enhance the natural color of coat, gives gloss and wetting, intensifies the natural waves of the hair and combats free radicals that damage the hair structure, preventing the hair breakage and its further loss.

THERAPEUTIC ACTION

Conditioner.

INGREDIENTS

Purified water, Propylene glycol, Behentrimonium Methosulfate/Cetyl Alcohol/Butylen glycol, Glycerin, Polyquaternium-37/ Polydecene Hydrogenated and Trideceth-6, Cetearyl Alcohol, Cetearyl Alcohol /Ceteareth-20, Bis-Isobutyl PEG/PPG-20/35/Amodimethicone Copolymer/ Cetyl Ethylhexanoate/ Polysorbate 80/Butylen glycol, Vitamin E, Fragrance, Disodium EDTA.

PROPERTIES

Maximum detangling effect. Cotton fragance.

USE INSTRUCTIONS

1. Apply on the back sufficient quantity of SIR DOG® CONDITIONER, on the moist and clean coat.
2. Massage gently until the balsam reaches the skin under the coat. Let act for 3 to 5 minutes.
3. Rinse with plenty of warm water until removing totally the balsam from the coat of your pet.
4. For obtaining a better result, it is recommended to bathe initially your dog with a shampoo of the SIR DOG® line.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

For external use.

CONDITION OF SALE

OTC product.

PRESENTATION

390 mL

PREPARED BY

Drag Pharma Laboratory.

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SIR DOG® ODOR CONTROL - SHAMPOO

SHAMPOO.

SHAMPOO NEUTRALIZADOR DE OLORES.

Technical Specification

SPECIES

Dogs.

Sir DOG ODOR CONTROL is specially formulated for neutralizing unpleasant odors of your pet's coat. Its composition allows to remove effectively the agents producing the bad odor in the canine skin. Also, it preserves the natural skin conditions, helping to keep a strong, soft and glossy hair.

DOSAGE FORM

Shampoo.

THERAPEUTIC ACTION

Shampoo neutralizes odors

PROPERTIES

Shampoo neutralizes odors with Piroctone Olamine.

USE INSTRUCTIONS

1. Wet your dog's coat completely with warm water.
2. Apply a sufficient quantity of Sir DOG ODOR CONTROL shampoo to reach the skin.
3. Massage until obtaining abundant lather, and let act for 3 to 5 minutes.
4. Rinse with plenty of warm water. For obtaining a better result, initially it is recommended to bathe your dog once a week for 4 weeks. Then, continue following the usual frequency of baths.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

For external use.

CONDITION OF SALE

OTC product.

PRESENTATION

390 mL

PREPARED BY

Drag Pharma Laboratory.

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SIR DOG® PERFUM LADY FLORAL WOOF

PERFUME.

EAU DE PERFUM.

Technical Specification

SPECIES

Dogs.

DOSAGE FORM

Eau de Parfum.

THERAPEUTIC ACTION

Eau de Parfum.

INGREDIENTS

Purified water, Cosmetic alcohol, Perfume, PEG-40 hydrogenated castor oil, Triclosan.

PROPERTIES

Sir DOG Lady is a perfume specially formulated for female dogs of any breed. An exquisite oriental Floral fragrance in accord with Rose and Apricot flower petals, carefully combined with notes of Lilac and Lily, with a warm background provided by Sandalwood, makes Sir DOG Lady the perfect fragrance for your pet. Sir DOG Lady has cosmetic alcohol, which is gentle on your pet's skin. The high quality of the base extracts and their concentration allow a prolonged permanence in the animal's fur.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

In order to not frighten your dog during the application, do not spray the product on the face or eyes.

CONSERVATION

Store in a cool, dry place protected from light, at no more than 30ºC.

CONDITION OF SALE

OTC product.

PRESENTATION

80 mL

PREPARED BY

Drag Pharma Laboratory.

AVAILABLE FOR SALE

COUNTRIES WHERE IT IS MARKETED

Imported and Distributed in Peru:
Representations Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604
Santiago de Surco Lima.Imported and Distributed in Peru:

SIR DOG® PERFUM LADY SWEETY LOVER

PERFUME.

EAU DE PERFUM.

Technical Specification

SPECIES

Dogs.

DOSAGE FORM

Eau de Parfum.

THERAPEUTIC ACTION

Eau de Perfum.

INGREDIENTS

Purified water, Cosmetic alcohol, Perfume, PEG-40 hydrogenated castor oil, Triclosan.

PROPERTIES

Sir DOG Lady Sweety Lover is a perfume specially formulated for female dogs of any breed. An exquisite feminine fragrance in accord with Berries of the Forest, Ripe Strawberries and Muguet Flower, carefully combined with notes of Freesia Flowers and Green Apple, with a warm background provided by Sandalwood and Musk. This balance of scents makes Sir DOG Lady Sweety Lover the perfect fragrance for your pet. Sir DOG Lady Sweety Lover has cosmetic alcohol, which is gentle on your pet's skin. The high quality of the base extracts and their concentration allow a prolonged permanence in the animal's fur.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a cool, dry place protected from light, at no more than 30 C.

CONDITION OF SALE

OTC Product.

PRESENTATION

80 mL

PREPARED BY

Drag Pharma Laboratory.

SIR DOG® PERFUM MUSK STRONGER LOVER

PERFUME.

EAU DE PERFUM.

Technical Specification

SPECIES

Dogs.

DOSAGE FORM

Eau de Parfum.

THERAPEUTIC ACTION

Eau de Perfum.

INGREDIENTS

Purified water, Cosmetic alcohol, Perfume, PEG-40 hydrogenated castor oil, Triclosan.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a cool, dry place protected from light, at no more than 30 C.

CONDITION OF SALE

OTC Product.

PRESENTATION

80 mL

PREPARED BY

Drag Pharma Laboratory.

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:
ZOOPHARMA
TEL: +(591)222-3357
1339 Díaz Romero Street, between Saavedra Avenue and Busch Avenue, La Paz.

Imported and Distributed in Peru:
Representations Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604
Santiago de Surco Lima.

SIR DOG® PERFUM MUSK WOODEN WOOF

PERFUME.

EAU DE PERFUM.

Technical Specification

SPECIES

Dogs.

DOSAGE FORM

Eau de Parfum.

THERAPEUTIC ACTION

Eau de Perfum.

PROPERTIES

Sir DOG Musk is a perfume specially formulated for dogs of any breed. An herbal and woody fragrance, with aromatic freshness delivered by Lavender and Rosemary, enhanced by Bitter Orange. With soft notes of sage and geranium, which linger alongside the warmth of cedar and vetiver.

This combination makes Sir DOG Musk the perfect fragrance for your pet.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a cool, dry place protected from light, at no more than 30 C.

CONDITION OF SALE

OTC Product.

PRESENTATION

80 mL

PREPARED BY

Drag Pharma Laboratory.

COUNTRIES WHERE IT IS MARKETED

Imported and Distributed in Peru:
Representations Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604
Santiago de Surco Lima.

SIR DOG® SHED CONTROL - SHAMPOO

SHAMPOO.

SHAMPOO PARA EL CONTROL DE LA CAÍDA DE PELO.

Technical Specification

SPECIES

Dogs.

SIR DOG® SHED CONTROL is specially formulated to prevent hair loss in your pet. Its composition, enriched with Vitamin E and Omega-3 and -6 Fatty acids, favors a healthy a glossy coat.

DOSAGE FORM

Shampoo.

THERAPEUTIC ACTION

Shampoo prevent the hair loss.

INGREDIENTS

Purified water, Lauryl ether sulfate sodium salt, Cocamidopropyl betaine, Cocamide DEA, Polysorbate 80, Propylene glycol, Sodium chloride, Soy oil, Vitamin E acetate, PEG-150 pentaerythrityl tetrastearate and PEG-6 caprylic/capric glycerides, Blue Colorant 1, Fragrance, Formaldehyde 37%, EDTA disodium salt, Citric acid anhydrous.

PROPERTIES

Shampoo with Vitamin E, Omega-3 and Omega-6.

USE INSTRUCTIONS

1. Wet your dog’s coat completely with warm water.
2. Apply SIR DOG® SHED CONTROL in a quantity enough to reach the skin.
3. Massage until obtaining abundant lather, and let act for 3 to 5 minutes.
4. Rinse with plenty of warm water.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

External use.

CONDITION OF SALE

OTC product.

PRESENTATION

390 mL

PREPARED BY

Drag Pharma Laboratory.

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SIR DOG® WHITE - SHAMPOO

SHAMPOO.

SHAMPOO PARA PERROS DE PELAJE BLANCO.

Technical Specification

SPECIES

Dogs.

SIR DOG® WHITE is specially formulated to enhance the white color of your pet's coat. Its composition, enriched with Optical Brightener and Aloe Vera Extract, favoring the condition of a healthy and glossy coat.

DOSAGE FORM

Shampoo.

THERAPEUTIC ACTION

Shampoo with Optical brightener and Aloe Vera extract.

INGREDIENTS

Purified water, Sodium Lauryl Ether Sulfate, Cocamidopropyl Betaine, Cocamide DEA, Glycol Distearate and Laureth-4, Propylenglycol, Sodium Chloride, Aloe Vera extract, Essence, Formalin 37%, Disodium EDTA , citric acid anhydrous, N,N-Dimethylformamide.

PROPERTIES

SIR DOG® WHITE is specially formulated to enhance the white color of your pet's coat. Coconut Fragrance.

USE INSTRUCTIONS

1. Wet all the coat of your pet using warm water.
2. Apply a sufficient quantity of SIR DOG® WHITE, for reaching the scalp.
3. Massage until obtaining abundant lather, and let act for 3 to 5 minutes.
4. Rinse with plenty of warm water.

WARNINGS

Mantener fuera del alcance de los niños.

CONDITION OF SALE

OTC product.

PRESENTATION

390 mL

PREPARED BY

Drag Pharma Laboratory.

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SKINDRAG® CERAMIDES - SHAMPOO

SHAMPOO CON CERAMIDAS PARA PELAJE DAÑADO Y OPACO.

SHAMPOO PREMIUM CON ESENCIA DE CRANBERRIES. REPARACIÓN & BRILLO.

Technical Specification

SPECIES

Dogs.

DESCRIPTION

SKINDRAG® Ceramides is a shampoo developed with Ceramides A2, a fatty acid that gives protective and repairing properties of the hair structure. It is hypoallergenic and its continuous use provides strength and brightness, restoring shine and softness to your pet coat.

DOSAGE FORM

Shampoo with Ceramides.

Damaged and dull coat.

THERAPEUTIC ACTION

Premium Dog Shampoo with Cranberries Essence. Repair & Shine.

INGREDIENTS

Lauryl Ether Sulfate Sodium Salt, Lauroyl Sarcosinate Sodium Salt, Sodium Chloride, Cocamide DEA, Glycerin, Ceramide A2, PEG-150 Pentaerythrityl Tetrastearate/PEG-6 Caprylic Glyceride, EDTA Disodium Salt, Glycol Distearate in Laureth-4, Cocamidopropyl Betaine, Fruits of the Forest Essence, Chloromethylisothiazolinone, Methylisothiazolinone, Purified water.

MODE OF APPLICATION

- Wet your dog’s coat completely with warm water.
- Apply a quantity of SKINDRAG® Ceramides enough to cover all coat.
- Massage until obtaining abundant lather.
- Rinse with plenty of warm water.

WARNINGS

Mantener fuera del alcance de los niños.

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SKINDRAG® OATS - SHAMPOO

SHAMPOO CON AVENA PARA PELAJE EXTRA SECO Y DELICADO.

SHAMPOO PREMIUM CON ESENCIA DE GROSELLAS, PARA HIDRATACIÓN & CUIDADO.

Technical Specification

SPECIES

Dogs.

DESCRIPTION

SKINDRAG® Oats is a shampoo will all the nourishing and dermocosmetic properties of Colloidal Oats. Its trace elements provides a protective coating on the skin, helping to delay the moisture evaporation from the skin, and thus prevent damaging the hydrophilic layer of animal skin. The oats particles absorb dirt and cell debris, preserving and caring the skin structure. On the other hand, it promotes the production of keratin, which adds strength and shine to the pet coat.

DOSAGE FORM

Premium Dog Shampoo. Extra Dry and Delicate Coat.

THERAPEUTIC ACTION

Premium Dog Shampoo.

INGREDIENTS

Lauryl Ether Sulfate Sodium Salt, Lauroyl Sarcosinate Sodium Salt, Sodium Chloride, Cocamide DEA, Glycerin, Colloidal Oats, PEG-150 Pentaerythrityl Tetrastearate/PEG-6 Caprylic Glyceride, EDTA Disodium Salt, Glycol Distearate in Laureth-4, Cocamidopropyl Betaine, Liquid Currant Essence, Chloromethylisothiazolinone, Methylisothiazolinone, Purified water.

MODE OF APPLICATION

- Wet your dog’s coat completely with warm water.
- Apply a quantity of SKINDRAG® Oats enough to cover all coat.
- Massage until obtaining abundant lather.
- Rinse with plenty of warm water.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Store in a fresh, dry place, out of direct light, at no more than 30 °C.

CONDITION OF SALE

OTC product.

PRESENTATION

250 mL

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SKINDRAG® VITAMIN E - CONDITIONER

ACONDICIONADOR PARA PELAJE LARGO O RESECO.

ACONDICIONADOR PREMIUM CON VITAMINA E, CON ESENCIA DE MIEL. SEDOSIDAD, BRILLO & PROTECCIÓN.

Technical Specification

SPECIES

Dogs.

DESCRIPTION

SKINDRAG® Conditioner has a formula rich in Vitamin E, which through its antioxidant effect, helps to reconstitute the internal matter of weakened fur, prevent color changes, restore the natural gloss of your pet coat, leaving it stronger and silkier.

DOSAGE FORM

Premium Dog Conditioner.

THERAPEUTIC ACTION

Premium Dog Conditioner with Vitamin E and Honey Essence. Silkiness, Gloss & Protection.

INGREDIENTS

Cetearyl Alcohol; Behentrimonium methosulfate; Cetyl Alcohol; Butylen glycol; Ceteareth 20; Bis-isobutyl Peg/PPG-20/35/ Amodimethicone Copolymer, Cetyl ethylhexanoate, Polysorbate 80; Polyquaternium-6; Vitamin E Oil; EDTA; Propylenglycol; Glycerin; Chloromethylisothiazolinone, Methylisothiazolinone; Honey essence; Purified water.

MODE OF APPLICATION

- Immediately after bathing your pet with SKINDRAG® shampoo, apply a sufficient quantity of SKINDRAG® Conditioner and massage, making sure that the product reaches all the coat.
- Rinse with plenty of warm water.

WARNINGS

Mantener fuera del alcance de los niños.

CONSERVATION

Keep in a fresh, dry place, out of the direct light, at no more than 30°C.

CONDITION OF SALE

OTC product.

PRESENTATION

250 mL

PREPARED BY

Drag Pharma Laboratory.

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SUAVIPET® TOPIC SOLUTION

SOLUCIÓN TÓPICA.

LOCIÓN DESENREDANTE.

Technical Specification

SPECIES

Dogs and cats.

It helps to detangle your pet coat. It provides silkiness and shine to coat with a pleasant coconut fragrance.For frequent use.

DOSAGE FORM

Topic solution.

THERAPEUTIC ACTION

Detangling lotion.

COMPOSITION

Deionized water, Amodimethicone, Dimethicone, Phenyl Trimethicone, fragrance and wetting agents.

PROPERTIES

Dogs and cats coat detangling spray lotion. SUAVIPET® is specially formulated with emollients and moisturizers to help detangling your pet's coat.

MODE OF APPLICATION

- Apply directly on the more tangled or knotted areas of coat, and then brush it until the knots are untied.
- SUAVIPET® provides softness and shine to coat, and also leaves an exquisite coconut fragrance.
- Use it regularly to prevent the coat tangles up again.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

For external use.

CONSERVATION

Store in a fresh, dry place, out of direct light, at no more than 30°C.

CONDITION OF SALE

OTC product.

PRESENTATION

200 mL

PREPARED BY

Drag Pharma Laboratory.

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SUPERPET® OMEGA ADULT DOG - ORAL SOLUTION

SOLUCIÓN ORAL.

SUPLEMENTO NUTRICIONAL DE ÁCIDOS GRASOS Y VITAMINA E PARA PERROS

Technical Specification

SPECIES

Adults dogs.

Superpet® Omega Adult Dog, is a mixture of oils of natural origin and Vitamin E, specially formulated for the maintenance of the health of your dog.
Its content in essential fatty acids Omega 6, favors growth in young animals, as well as the development and maintenance of healthy skin and coat for your pet. Omega 3 fatty acids provide a natural source of protection for the functioning of the nervous, cardiovascular and immune systems.

Due to its antioxidant activity, it helps to slow down the aging processes.

- Omega 6 essential fatty acids help maintain healthy skin and healthy coat throughout your pet's life.
- The optimal ratio of Omega 6 / Omega 3 fatty acids favors your dog's metabolism.

DOSAGE FORM

Oral Solution.

THERAPEUTIC ACTION

Nutritional Supplement of Fatty Acids and Vitamin E for dogs

INGREDIENTS

Sunflower Oil, Borage Oil, Salmon Oil, DL-Alpha-Tocopherol Acetate (Vitamin E), authorized flavorings.

NUTRITIONAL CONTENT

Each 1 mL contains:
Linoleic Acid (LA).....439.8 mg
Gamma-Linolenic Acid (GLA).....9.96 mg
Eicosapentanoic Acid (EPA).....21.06 mg
Docosahexanoic Acid (DHA).....28.91 mg
Vitamin E (as Acetate).....46 IU
Omega-6 : Omega-3 ratio = 8.6 : 1

PROPERTIES

ROUTE OD ADMINISTRATION AND DOSAGE

3 mL for every 10 Kg of weight (3 pumps a day).

Administer mixed with food or directly orally.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

Meat flavor.

It does not constitute a complete food.
Exclusive use in animal feed

CONSERVATION

Store in a cool, dry place and away from light, at no more than 30ºC.

CONDITION OF SALE

OTC product (non-prescription)

PRESENTATION

Dosing bottle with 125 mL

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Chile: Reg. LENAA N°: RM 03-008N
Panamá: Reg. Nº RF-4187-19
Bolivia: Reg. SENASAG PUV-A-N° 005510/13
El Salvador: AL2014112627
Costa Rica: Lic. DAA-MAG 579-012
Perú: Reg. SENASA A.16.15.I.0312

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

Distribution in El Salvador: Rafael Alfredo Alfaro Castillo.
8a C. Pte. Pje. Moreno N ° 112, Col. Flor Blanca.
San Salvador, El Salvador.

Imported and distributed in Costa Rica by:
Proventas de Cartago S.R.L. 100 meters east Hogares Crea, San Blas. Carthage.
Tel: 2591 4624 Fax: 2591 5339

Imported and Distributed in Peru by: Representaciones Durand SAC.
Av. Manuel Olguín N ° 501 Office N ° 604 Santiago de Surco Lima.

SUPERPET® OMEGA CAT - ORAL SOLUTION

SOLUCIÓN ORAL.

SUPLEMENTO NUTRICIONAL DE ÁCIDOS GRASOS Y VITAMINA E.

Technical Specification

SPECIES

Cats.

DOSAGE FORM

Oral solution.

THERAPEUTIC ACTION

Nutritional Supplement of Fatty Acids and Vitamin E.

INGREDIENTS

Sunflower oil, Borage oil, Salmon oil, DL-Alpha Tocopheryl Acetate (Vitamin E), flavors.

NUTRITIONAL CONTENT

Each 1 mL contains:

Linoleic Acid (LA).....439.8 mg

Gamma-Linolenic Acid (GLA).....9.96 mg

Eicosapentanoic Acid (EPA).....21.06 mg

Docosahexanoic Acid (DHA).....28.91 mg

Vitamin E (as Acetate).....46 IU

Omega-6 : Omega-3 ratio = 8.6 : 1.

PROPERTIES

SUPERPET® OMEGA 6 : 3 is a mixture of natural oils and Vitamin E, specially formulated to preserve your cat health. Its content in essential Omega-6 fatty acids favors the growing in young animals, as well the development and preservation of a healthy skin and coat for your pet. The Omega-3 fatty acids provide a natural source of protection for the functioning of the nervous, cardiovascular and immune systems. Because its antioxidant action, it helps to delay the inherent processes of aging. The essential Omega-6 fatty acids help to recover a healthy skin after a dermatitis, improving also the condition of coat. The optimal ratio of omega 6/omega 3 fatty acids favors the metabolism in your cat.

ROUTE OD ADMINISTRATION AND DOSAGE

- Growing cat: 1 Dose (1 pumping / day)
- Adult cat (4 kg): 2 Doses (2 pumpings / day)

Administer mixed with food or directly by oral route.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

- Crab flavor.
- It is not a complete food.

CONSERVATION

Store in a fresh, dry place, out of direct light, at no more than 30°C.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

125 mL

PREPARED BY

Drag Pharma Laboratory.

Imported and distributed by: Bolivia Interagro S.A. Reg. SENASAG PUV-AN° 005518/13. Tech. Resp. Dr. Fernando Vargas (Matr. 983)

RECORDS

- SAG Inscription #: RM03-010
- Panama: Reg. No. RF-4184-08

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SUPERPET® OMEGA PUPPY - ORAL SOLUTION

SOLUCIÓN ORAL.

SUPLEMENTO NUTRICIONAL DE ÁCIDOS GRASOS Y VITAMINAS.

Technical Specification

SPECIES

Dogs (puppies).

DOSAGE FORM

Oral solution.

THERAPEUTIC ACTION

Nutritional Supplement of Fatty Acids and Vitamins.

INGREDIENTS

Sunflower oil, Borage oil, Linseed oil, DL-Alpha Tocopheryl Acetate, Vitamin A palmitate, Vitamin D, Authorized flavors.

NUTRITIONAL CONTENT

Each 1 mL contains:

Linoleic Acid (LA) .....350.0 mg (min.)

Gamma-Linolenic Acid (GLA) .....8.5 mg (min.)

Alpha-Linolenic Acid (ALA).....60.0 mg (min.)

Oleic Acid (OA).....180.0 mg (min.)

Vitamin E.....46 IU

Vitamin A.....200 IU

Vitamin D.....20 IU

Omega 6 : Omega 3 ratio = 10:1 to 5: 1

PROPERTIES

Superpet® omega puppy is a nutritional supplement that provides a balanced supply of essential omega 6 and omega 3 fatty acids, necessary during the first months of life and during the vaccination period.

Its content of essential fatty acids and vitamins favors an excellent health of the skin and coat of a growing puppy. Superpet® Omega Puppy allows to reduce the continuous hair loss, the excessive scratching, and to avoid the appearance of a dull coat and squamous skin, very common in this step of development, since its composition allows to cover the increasing nutritional needs of the puppy.

The vitamins present in Superpet® Omega Puppy contribute to the health care of your puppy, because of its specific functionalities for the body wellness.

ROUTE OD ADMINISTRATION AND DOSAGE

1 ml for each 1 kg of body weight (1 pumping / kg daily)

Administer mixed with food or directly by oral route.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

It is not a complete food.

CONSERVATION

Store in a fresh, dry place, out of direct light, at no more than 30°C.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

125 mL

PREPARED BY

Drag Pharma Laboratory.

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SUPERPET® OMEGA SENIOR - ORAL SOLUTION

SOLUCIÓN ORAL.

SUPLEMENTO NUTRICIONAL DE ÁCIDOS GRASOS, VITAMINA E Y LECITINA.

Technical Specification

SPECIES

Dogs (senior).

Superpet® Omega Senior is a nutritional supplement that provides a balanced supply of essential fatty acids, vitamin E and lecithin for older dogs. Its content in essential omega-6 and omega-3 fatty acids favors an excellent skin and coat health for your dog. The Omega-3 fatty acids provide a natural source of protection for the functioning of the nervous, cardiovascular and immune systems. Its content in soy lecithin is an excellent supply of phospholipids: phosphatidylcholine, phosphatidylethanolamine and phosphatidylinositol, which allow to preserve the health of cell membranes, supporting the brain function and preventing a premature cell aging. Its contribution in Vitamin E neutralizes the free radicals generated by the aging, preventing the cell damage and delaying the cognitive impairment in older dogs.

DOSAGE FORM

Oral Solution.

THERAPEUTIC ACTION

Nutritional supplement of fatty acids, vitamin E and lecithin.

INGREDIENTS

Sunflower oil, Salmon oil, Borage oil, Vitamin E Acetate, Soy Lecithin, authorized flavors.

NUTRITIONAL CONTENT

Each 1 mL contains:

Linoleic Acid (LA).....320.0 mg (min.)

Gamma-Linolenic Acid (GLA).....8.0 mg (min.)

Eicosapentanoic Acid (EPA).....12.0 mg (min.)

Docosahexanoic Acid (DHA).....10.0 mg (min.)

Oleic Acid (OA).....120.0 mg (min.)

Lecithin.....10 mg (min.)

Vitamin E (as Acetate).....46 IU

Omega-6 : Omega-3 ratio = 10:1 to 5:1

PROPERTIES

It favors the increase of defenses, revitalizes the brain function, and improves the skin and coat health of your pet.

ROUTE OD ADMINISTRATION AND DOSAGE

3 ml for each 20 kg of body weight (3 pumpings / 20 kg daily).

Administer mixed with food or directly by oral route.

WARNINGS

Mantener fuera del alcance de los niños.

OBSERVATIONS

It is not a complete food.

CONSERVATION

Store in a fresh, dry place, out of direct light, at no more than 30°C.

CONDITION OF SALE

OTC product (non-prescription).

PRESENTATION

125 mL

PREPARED BY

Drag Pharma Laboratory.

Imported and distributed by: Bolivia Interagro S.A. Reg. SENASAG PUV-AN° 005512/13. Tech. Resp. Dr. Fernando Vargas (Matr. 983)

RECORDS

SAG Inscription #: RM03-010

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

SUPOLEN® ORAL TABLET

COMPRIMIDOS ORALES.

FASCIOLICIDA INTERNO DE AMPLIO ESPECTRO EN DOSIS ÚNICA.

Technical Specification

SPECIES

Cattle, sheep and goats.

Single dose broad spectrum internal fasciolicide. Effective against immature, juvenile and adult forms of *Fasciola hepatica*.

DOSAGE FORM

Oral tablet

THERAPEUTIC ACTION

Single dose broad spectrum internal fasciolicide.

COMPOSITION

Each tablet contains:
Triclabendazole ..... 1.2 g
Excipients q.s.p ..... 1 tablet

INDICATIONS

Do not use in animals with known hypersensitivity to Triclabendazole.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration way: Oral.

Dose of the active principle:
- Bovines: 12 mg / Kg of body weight.
- Sheep and Goats: 10 mg / Kg of body weight.

Product dosage:
- Cattle: 1 tablet per 100 Kg of weight in a single dose.
- Sheep and goats: ½ tablet for every 60 Kg of weight in a single dose.

DRUG INTERACTIONS

Do not use concomitantly with other pharmaceutical products.

CONTRAINDICATIONS

Do not use in animals with known hypersensitivity to Triclabendazole.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Avoid contact with eyes and skin. In case of contact, wash immediately with plenty of water.
Wash your hands after handling the product.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Meat: 28 days.

Milk: Do not use the milk of treated animals for human consumption.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 ° and 30 ° C, protected from light.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Pot with 50 tablets.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Reg. SAG N° 592

TENIMOX® ORAL PASTE

PASTA ORAL.

ANTIPARASITARIO INTERNO DE AMPLIO ESPECTRO CON DORAMECTINA EXCLUSIVO PARA CABALLOS

Technical Specification

SPECIES

Horses

Tenimox® is a broad spectrum antiparasitic, which thanks to its composition based on Doramectin and Praziquantel, is effective against nematodes, such as small and large strongyles, Parascaris equorum and Oxyuris equi. It is also effective against cestodes such as Anoplocephala perfoliata and obligatory myiases produced by parasites such as Gasterophilus spp. Tenimox® is effective for at least 80 days after administration.

DOSAGE FORM

Oral Paste

THERAPEUTIC ACTION

Broad spectrum internal antiparasitic with DORAMECTIN exclusive for horses

COMPOSITION

Each 100 g of product contains:
Doramectin ..... 1.2 g
Praziquantel ..... 9.0 g
Excipients q.s.p ..... 100.0 g

INDICATIONS

- Do not use in animals with liver and / or kidney damage.
- Do not use in foals under 5 months.
- Do not use on animals in poor condition or in stressful situations.
- Do not use in pregnant females.

CONTRAINDICATIONS

- Do not use in animals with liver and / or kidney damage.
- Do not use in foals under 5 months.
- Do not use on animals in poor condition or in stressful situations.
- Do not use in pregnant females.

WARNINGS

- Lavar las manos después de administrar el producto a los animales.
- Mantener fuera del alcance de los niños.

GUARD PERIOD

Do not administer to horses whose meat is intended for human consumption.

OBSERVATIONS

Cherry flavor.

CONSERVATION

Keep in a cool and dry place, between 2 and 30 ° C, protected from light.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

Dosing syringe with 10 g

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 2054-B
Bolivia: Reg. SENASAG PUV-F N° 007262/16
Uruguay: Reg. MGAP N° 2017A00409

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:
AGROGUARANI SRL
TEL: + (591) 314-1401
Santa Cruz de la Sierra, Bolivia

Importer in Uruguay:
VIVAFIL S.A.
RIO NEGRO 1107 Montevideo - Uruguay, TEL 29001112
grupotecnovet@gmail.com
Technical Director: DMTV Diego Cuadrado.

TERIL® ORAL SUSPENSION

SUSPENSIÓN ORAL.

ANTIESPUMANTE Y ANTITIMPANIZANTE RUMINAL.

Technical Specification

SPECIES

Cattle, goats, sheep and horses.

Bloat or weathering (filling).
Gastric or ruminal overload, gaseous colic and in general in all cases in which the normal process of rumination is disturbed.

DOSAGE FORM

Oral suspension

THERAPEUTIC ACTION

Antifoam and ruminal antimpanizer

COMPOSITION

Each 100 mL contains:
Simethicone 30% emulsion ..... 5 g
Excipients q.s.p ..... 100 mL

INDICATIONS

Do not administer in animals with hypersensitivity to the active principle.

ROUTE OD ADMINISTRATION AND DOSAGE

- Adult cattle: 100 - 250 mL directly intraruminally, orally, pure or mixed with water.
- Small ruminants: 50 - 100 mL directly intraruminally, orally, pure or mixed with water.
- Horses: 100 - 200 mL, orally alone or dissolved in water.

CONTRAINDICATIONS

Do not administer in animals with hypersensitivity to the active principle.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Avoid contact with eyes and skin. In case of contact with the eyes, wash immediately with plenty of water for 15 minutes. If there is contact with the skin, wash with soap and water. If the discomfort persists, consult a doctor.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Meat and milk: 24 hours.

OBSERVATIONS

SHAKE BEFORE USING

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of natural water. Dispose of this product waste carefully with household waste.

CONSERVATION

Keep in a cool and dry place, at room temperature between 15 and 30 ° C, protected from light. Use immediately once opened and discard the excess product.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

500 mL bottle and 5 liter drum.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

- Chile: Reg. SAG N° 408

TIDY® CATS - DRY SHAMPOO

SHAMPOO SECO.

ASEO EN SECO.

Technical Specification

SPECIES

Cats.

Dry shampoo for cats.

Tidy® is specially formulated for dry cleaning.
Tidy®, in addition to the principles for grooming the coat, contains fine essences to make the presence of your pet more pleasant.

DOSAGE FORM

Dry shampoo.

THERAPEUTIC ACTION

Perfumed dry shampoo.

INGREDIENTS

Corn Starch, Sodium Chloride, Talc, Magnesium Carbonate, Carboxymethylcellulose, Borax, Authorized fragrance.

MODE OF APPLICATION

- Do not get your pet wet.
- Sprinkle in a sufficient amount of Tidy®, scrub vigorously with your hands and finish by brushing gently.

OBSERVATIONS

For external use.

CONSERVATION

Store in a cool, dry place, protected from light, at no more than 30ºC.

PRESENTATION

100 g talcum powder container

PREPARED BY

Drag Pharma Laboratory.

RECORDS

Costa Rica: Reg. MAG Nº CL-4-45-13-6018
Bolivia: Reg. SENASAG PUV N° 008272/18

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:

ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

TIDY® DOGS - DRY SHAMPOO

SHAMPOO SECO.

ASEO EN SECO.

Technical Specification

SPECIES

Dogs.

Tidy® is specially formulated for dry cleaning.
Tidy®, in addition to the principles for grooming the coat, contains fine essences to make the presence of your pet more pleasant.

DOSAGE FORM

Dry shampoo.

THERAPEUTIC ACTION

Perfumed dry shampoo.

INGREDIENTS

Corn Starch, Sodium Chloride, Talc, Magnesium Carbonate, Carboxymethylcellulose, Borax, Authorized fragrance.

MODE OF APPLICATION

- Do not get your pet wet.
- Sprinkle in a sufficient amount of Tidy®, scrub vigorously with your hands and finish by brushing gently.

OBSERVATIONS

External use.

CONSERVATION

Store in a cool, dry place, protected from light, at no more than 30ºC.

PRESENTATION

100 g talcum powder container

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Costa Rica: Reg. MAG CL4-45-13-6221
Bolivia: Reg. SENASAG PUV N° 008266/18

AVAILABLE FOR SALE

Importado y distribuido en Bolivia por:

ZOOFARMA
TEL: +(591)222-3357
Calle Díaz Romero 1339, entre avenida Saavedra y Avenida Busch, La Paz.

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:
AGROGUARANI SRL
TEL: + (591) 314-1401
Santa Cruz de la Sierra, Bolivia

TONIMAG® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

MAGNESIO INYECTABLE.

Technical Specification

SPECIES

Bovines and sheep.

In the prevention and treatment of clinical and subclinical hypomagnesemia. Metabolic imbalances due to nutritional deficiencies during lactation, caused by the consumption of fresh pastures or rich in cereals or in animals that remain outdoors in winter exposed to sudden climatic changes (rains, frosts).

DOSAGE FORM

Injectable solution

THERAPEUTIC ACTION

Injectable magnesium

COMPOSITION

Each 100 mL of solution for injection contains:
Magnesium Sulfate x 7 H_2O ..... 46.6 g
(Equivalent to 4.6 g of Magnesium)
Magnesium Glycerophosphate. ..... 4.64 g
(Equivalent to 0.58 g of Magnesium)
Potassium Chloride ..... 3 g
(Equivalent to 1.57 g of Potassium)
Excipients q.s.p... ..... 100 mL

INDICATIONS

- Do not administer in animals with heart and / or kidney failure.
- Do not administer in animals with hyperkalemia or hypermagnesemia.
- Do not administer in animals with hemolytic disorders, untreated Addison's disease and acute dehydration.

DRUG INTERACTIONS

It is not recommended to use concomitantly with other pharmaceutical products.

CONTRAINDICATIONS

- Do not administer in animals with heart and / or kidney failure.
- Do not administer in animals with hyperkalemia or hypermagnesemia.
- Do not administer in animals with hemolytic disorders, untreated Addison's disease and acute dehydration.

PRECAUTIONS

- In case of observing a decrease in the respiratory and cardiac frequency or alterations in the respiratory depth, suspend the administration until the normal parameters are reestablished.
- Observe and monitor the animal at all times.

SPECIAL PRECAUTIONS FOR USE

- In case of observing a decrease in the respiratory and cardiac frequency or alterations in the respiratory depth, suspend the administration until the normal parameters are reestablished.
- Observe and monitor the animal at all times.

SPECIAL PRECAUTIONS FOR THE OPERATOR

Handle the product with care. In case of accidental self-injection, immediate medical attention should be sought.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

The use of Magnesium and Potassium in high doses can cause:

- At the CNS level: Somnolence and neuromuscular weakness.
- At the Cardiorespiratory level: Bradycardia, hypotension, hypocalcemia, respiratory depression and increases in Q-T intervals on the electrocardiogram. Eventually very high doses can cause cardiac arrest.
- Clinical signs can be exacerbated by concomitant hypocalcemia, hyponatremia, or acidosis.

GUARD PERIOD

Meat: 0 days.

Milk: 0 days.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard the remains of unused product in its original container, well closed. Do not throw the empty container or with product remains in the ground, rivers or streams of water, or reuse the container. Dispose of this product waste carefully with household waste.

CONSERVATION

Store at room temperature between 2 and 30 ° C, protected from light. Once the container is opened, use within 1 month and store between 15 and 30 ° C, protected from light. Discard the unused product after that period of time.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

100 mL vial

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

- Chile: Reg. SAG N° 1465
- Costa Rica: Reg. N° MAG CL4-34-01-3672

TOPFENICOL® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANTIMICROBIANO PARA BOVINOS Y CERDOS.

Technical Specification

SPECIES

Bovines and pigs.

In cattle, it is indicated in bovine respiratory disease, in which susceptible strains of Pasteurella haemolitica, Pasteurella multocida, Haemophilus somnus, Actinomyces pyogenes are associated. Its therapeutic action is also recommended in infectious keratoconjunctivitis caused by Moraxella bovis.

In pigs, its use is related to respiratory disease. The strains causing the syndrome that are described as susceptible are Pasteurella multocida, *Mycoplasma hyopneumoniae, Salmonella cholerasuis, Mycoplasma hyorhinus, Bordetella bronchiseptica, Streptococcus suis.*

DOSAGE FORM

Injectable solution

THERAPEUTIC ACTION

Antimicrobial for cattle and pigs.

COMPOSITION

Each 1 mL of product contains:
Florfenicol ..... 300 mg
Excipients q.s.p ..... 1 mL

PROPERTIES

Florfenicol is a long-acting injectable antibiotic with action on a wide variety of aerobic and anaerobic Gram (+) and Gram (-) bacteria, isolated from cattle and pigs.

INDICATIONS

- Do not administer to animals that have previously shown hypersensitivity to chloramphenicol or florfenicol.
- Do not use in cattle intended for encasing.
- Do not use on bristles intended for setting.
- Do not use in pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Dose of the active ingredient:

- Cattle: 20 mg / Kg of live weight, intramuscularly.
- Pigs: 15 mg / Kg of live weight, intramuscularly.

Repeat a second dose 48 hours after starting treatment.

Product dosage:

- Cattle: 6.7 mL / 100 Kg of weight, intramuscularly.
- Pigs: 0.5 mL / 10 Kg of live weight, intramuscularly.

Repeat a second dose 48 hours after starting treatment.

CONTRAINDICATIONS

- Do not administer to animals that have previously shown hypersensitivity to chloramphenicol or florfenicol.
- Do not use in cattle intended for encasing.
- Do not use on bristles intended for setting.
- Do not use in pregnant or lactating females.

PRECAUTIONS

If signs of respiratory disease persist 48 hours after the second dose, reevaluate the diagnosis.

SPECIAL PRECAUTIONS FOR USE

If signs of respiratory disease persist 48 hours after the second dose, reevaluate the diagnosis.

ADVERSE EFFECTS

In cattle, a temporary drop in food and water consumption may occur at the start of therapy.
In pigs, transient perianal inflammation may occur at the start of treatment.

OBSERVATIONS

Employment during pregnancy and lactation:
Do not use in animals intended for reproduction, or in pregnant or lactating females. Efficacy period: 36 months

Efficacy period
36 months

Disposal of waste material:
Discard any unused product remains in its original container. Do not throw the empty container or with the rest of the product in rivers, lakes or streams of water. Dispose of the remains of this product with caution together with household waste.

CONSERVATION

Store between 15 and 30 ° C, in a cool and dry environment, protected from light and in its original container.

CONDITION OF SALE

To be supply only on veterinary prescription.

PRESENTATION

100 mL vial

PREPARED BY

Laboratorio Drag Pharma Chile Invetec.S.A.

RECORDS

Chile: Reg. SAG N° 2022
Bolivia: Reg. SENASAG PUV-F nº 007492/16
Perú: Registro SENASA F.03.01.I.1844

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Bolivia by:
ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

Imported and Distributed in Peru by Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

TOPFLAM® TOPICAL GEL

GEL TÓPICO.

ANTIINFLAMATORIO NO ESTEROIDAL

Technical Specification

SPECIES

Horses.

Topflam® is indicated for the control of pain and inflammation associated with the musculoskeletal system.

DOSAGE FORM

Topical Gel

THERAPEUTIC ACTION

Anti-inflammatory

COMPOSITION

Each 100 g of product contains:
Diclofenac Sodium ..... 1 g
Excipients q.s.p ..... 100 g

INDICATIONS

- Do not administer in animals sensitive to the drug.
- Do not use in conjunction with NSAIDs.
- Do not administer during pregnancy or lactation.
- Do not administer to foals under one year of age.

ROUTE OD ADMINISTRATION AND DOSAGE

Administer topically 7.3 g of gel or a 13 cm band of gel, twice a day for 5 days, equivalent to 73 mg of Diclofenac Sodium in each application. Rub the joint to be treated with the product until it is completely absorbed.

DRUG INTERACTIONS

The use of other topical drugs should be avoided as they can inhibit the absorption of Topflam® and cause irritation in the area of application. The concomitant administration of other NSAIDs by the systemic route has not been studied.

CONTRAINDICATIONS

- Do not administer in animals sensitive to the drug.
- Do not use in conjunction with NSAIDs.
- Do not administer during pregnancy or lactation.
- Do not administer to foals under one year of age.

PRECAUTIONS

Exceeding the recommended dose or treating several joints at the same time in the same animal, can increase the plasma concentrations of Diclofenac. Keep out of reach of children and pets.

SPECIAL PRECAUTIONS FOR USE

Exceeding the recommended dose or treating several joints at the same time in the same animal, can increase the plasma concentrations of Diclofenac. Keep out of reach of children and pets.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Avoid contact with the skin, wear gloves to apply the product. In the case of contact, wash with soap and water.
- Avoid accidental contact with the eyes. In the case of contact, rinse with plenty of water and consult a doctor immediately.
- Do not handle by people who are hypersensitive to Diclofenac Sodium.
- In case of accidental ingestion, go immediately to a medical center and show the product label.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Do not administer to horses whose meat is intended for human consumption.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Keep in a cool and dry place, at room temperature between 15° and 30° C, protected from light. Once the container is opened, use within 12 weeks. Discard unused product after that time period.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

120 g tube

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 2042
Perú: Reg. SENASA F.C4.75.I.0010

COUNTRIES WHERE IT IS MARKETED

Imported and Distributed in Peru by Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

TRANSIMED® OTIC AND TOPICAL SUSPENSION

SUSPENSIÓN ÓTICA Y TÓPICA.

ANTIBACTERIANO. ANTIMICÓTICO. ANTIINFLAMATORIO ESTEROIDAL.

Technical Specification

SPECIES

Dogs and cats.

Transimed® is indicated for the treatment of external otitis and dermatitis caused by fungi, yeasts and Gram (-) bacteria.

DOSAGE FORM

Otic and topical suspension.

THERAPEUTIC ACTION

Antibacterial, antifungal and steroidal anti-inflammatory

COMPOSITION

Each 1 mL contains:
Miconazole Nitrate ..... 23.0 mg
Prednisolone Acetate ..... 5.0 mg
Polymyxin B Sulfate ..... 5,000 IU
Excipients q.s.p ..... 1 mL

INDICATIONS

- Do not use in animals with a perforated eardrum due to the ototoxic effect of Polymyxin B.
- Do not use in pregnant or lactating females.

ROUTE OD ADMINISTRATION AND DOSAGE

Route of administration: Topical.

Dosage and directions for use:
In otitis:
- Clean the external auditory canal, with a ceruminolytic and antiseptic product.
- Instill 5 drops of Transimed® in the external auditory canal 2 to 3 times a day, for up to 15 days.

In dermatitis:
- Cut the hair in the affected area and clean it.
- Spread several drops of Transimed® on the area to be treated, 2 to 3 times a day for up to 10 days.

DRUG INTERACTIONS

- Polymyxin B has interactions with aminoglycoside antibiotics, Bacitracin, Quinidine, Quinine.
- Sodium Citrate enhances the nephrotoxic and neurotoxic potential of Polymyxin B.
- Succinylcholine and Tubocurarine can prolong the neuromuscular blockade of respiratory paralysis associated with the use of Polymyxin B.

CONTRAINDICATIONS

- Do not use in animals with a perforated eardrum due to the ototoxic effect of Polymyxin B.
- Do not use in pregnant or lactating females.

PRECAUTIONS

In case of prolonged therapy (more than 7 days) it should be gradually suspended in successive days according to what is indicated by the treating Veterinarian.

SPECIAL PRECAUTIONS FOR USE

In case of prolonged therapy (more than 7 days) it should be gradually suspended in successive days according to what is indicated by the treating Veterinarian.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Do not handle by people who are hypersensitive to glucocorticoids.
- In case of contact with skin, eyes or mucous membranes, wash immediately with plenty of water.
- Do not handle by pregnant women.
- In case of ingestion, go immediately to a medical center and show the product label.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

- The use of Transimed® for more than 7 days could delay wound healing.
- The use of Transimed® may be associated with the development of deafness or partial hearing loss in a small number of sensitive (geriatric) dogs. The hearing deficit is usually temporary. If vestibular or hearing dysfunctions are detected during the course of treatment, therapy should be discontinued and the external auditory canal should be flushed.

OBSERVATIONS

Shake before using.

Special precautions for disposal of unused product or waste material:
Empty containers can be disposed of as household waste, without any special precautions.
Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store at room temperature, between 15 and 30 ° C. Once opened use the product within 3 weeks.
Discard the unused product after that period of time.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

15 mL bottle with cannula.

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG Nº 0526-B
El Salvador: VE2015105122

COUNTRIES WHERE IT IS MARKETED

Distribution in El Salvador:
Rafael Alfredo Alfaro Castillo.
8th C. Pte. Pje. Moreno N ° 112, Col. Flor Blanca.
San Salvador, El Salvador.

Imported and distributed in Bolivia by:
ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

TRIAMCOL® INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

ANALGÉSICO.

Technical Specification

SPECIES

Horses.

Triamcol®, Tramadol Hydrochloride 5%, Injectable solution, is indicated as an analgesic in the management of acute pain of low to severe intensity that accompanies colic in horses.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Analgesic

COMPOSITION

Each 1 mL of product contains:
Tramadol Hydrochloride ..... 50 mg
(Equivalent to 43.9 mg of Tramadol base)
Excipients q.s.p ..... 1 mL

PROPERTIES

Tramadol Hydrochloride is a synthetic opioid agonist that acts primarily through binding to μ receptors, in addition to inhibiting the reuptake of the monoamines serotonin and norepinephrine. It is from the two mechanisms mentioned above that the sensation of pain is inhibited at the central level, causing a powerful analgesia (antinociception) and a moderate anxiolytic effect.

INDICATIONS

- Do not administer to horses with known hypersensitivity to Tramadol.
- Do not administer to pregnant or lactating females.
- Do not administer in conjunction with Monoamine Oxidase Inhibitors (MAOIs) such as Selegiline and Isoniazid; Selective Serotonin Reuptake Inhibitors (SSRIs) such as Fluoxetine and Paroxetine; SSRIs and norepinephrine reuptake inhibitors such as Clomipramine; Serotonin antagonists such as Cyproheptadine; Noradrenaline antagonists such as Prazosin; Noradrenaline agonists such as Clonidine.

ROUTE OD ADMINISTRATION AND DOSAGE

Dose of the active principle:
2 mg of Tramadol Hydrochloride / Kg of body weight.

Product dosage:
4 mL / 100 Kg of body weight.

Routes and forms of administration: Intravenous administration.
The administration of the product should be slow (in 5 to 10 minutes) to minimize possible side effects.

CONTRAINDICATIONS

- Do not administer to horses with known hypersensitivity to Tramadol.
- Do not administer to pregnant or lactating females.
- Do not administer in conjunction with Monoamine Oxidase Inhibitors (MAOIs) such as Selegiline and Isoniazid; Selective Serotonin Reuptake Inhibitors (SSRIs) such as Fluoxetine and Paroxetine; SSRIs and norepinephrine reuptake inhibitors such as Clomipramine; Serotonin antagonists such as Cyproheptadine; Noradrenaline antagonists such as Prazosin; Noradrenaline agonists such as Clonidine.

SPECIAL PRECAUTIONS FOR THE OPERATOR

During the handling and application of the product, the use of gloves is recommended. In the case of accidental ingestion, contact or injection it is recommended:

- **First aid:** Persons who develop serious hypersensitivity (anaphylactic) reactions should receive immediate medical attention. If person is not breathing, give artificial respiration. If breathing is difficult, give oxygen. Get medical attention.
- **Risk of contact with the skin:** In the event of contact with the skin, it is recommended to wash immediately with plenty of water and soap. In the event of any type of irritation, it is recommended to see a doctor.
- **Risk of contact with the eyes:** In the event of contact with the eyes, it is recommended to wash immediately with plenty of water for approximately 15 minutes. In the event of any type of irritation, it is recommended to see a doctor.
- **Ingestion risk:** In the case of accidental ingestion, it is recommended to rinse the mouth with plenty of water. Do not induce vomiting unless directed by a doctor. It is recommended to go to a healthcare center promptly.

WARNINGS

Mantener fuera del alcance de los niños.

SIDE EFFECTS

In horses, nausea, salivation and tremors may be observed during intravenous administration of Tramadol Hydrochloride. These effects disappear if the administration by this route is done slowly (in 5 to 10 minutes). After intravenous administration, an increase in systolic blood pressure and tachycardia could occur in more sensitive individuals, which should remit within 30 to 60 minutes after administration.

GUARD PERIOD

Do not administer to horses whose meat is intended for human consumption.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Any unused medicine or the waste derived from it must be disposed of in an environmentally safe way and in accordance with local regulations. Empty containers or containers with the product should not be disposed of on the ground or in water courses.
These materials must be disposed of by authorized companies to perform said service safely.

CONSERVATION

Keep in a dry place, at a temperature between 2 and 30º C, protected from light.

CONDITION OF SALE

Sale with retained Veterinary Medical prescription.

PRESENTATION

50 mL vial

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 2169

Perú: Reg. SENASA F.G4.01.I.0001

Bolivia: Reg. SENASAG PUV- Nº 10000/21

AVAILABLE FOR SALE

Imported and distributed in Bolivia by:
ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

TRIPLE- INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

VITAMINAS B1 - B6 - B12.

Technical Specification

SPECIES

Cattle, sheep, goats, horses, pigs, dogs and cats.

Indicated in deficiency states of vitamins B_1, B_6 and B_{12}. As an adjuvant in the treatment of diseases with vitamin B_1, B_6 and B_{12} deficiencies.

DOSAGE FORM

Injectable solution

THERAPEUTIC ACTION

B1 - B6 - B12 Vitamins

COMPOSITION

Each 5 mL of injectable solution contains:
Thiamine Hydrochloride (Vitamin B1) ..... 220.0 mg
Pyridoxine Hydrochloride (Vitamin B6) ..... 220.0 mg
Cyanocobalamin (VItamine B12) ..... 21 mg
Excipients csp .... 5 mL

ROUTE OD ADMINISTRATION AND DOSAGE

Route of administration: Intramuscular.

- **Cattle and horses:** 5 to 10 mL, per day, in a single dose.
- **Pigs, sheep, goats, calves and foals:** 5 mL, per day, in a single dose.
- **Dogs and cats:** 1 to 2 mL, a day, in a single dose.

Repeat the dosage if necessary.

DRUG INTERACTIONS

B vitamins should not be mixed with other drugs.
Vitamin B in general is incompatible with: Chloramphenicol, Chlorpromazine Hydrochloride, Methylprednisolone, intravenous infusions of Sodium Bicarbonate.

WARNINGS

- Mantener fuera del alcance de los niños.
- No usar en animales hipersensibles a Tiamina administrada vía parenteral, por riesgo a shock anafiláctico.

GUARD PERIOD

0 days

OBSERVATIONS

Fractional sale is prohibited.

CONSERVATION

- Store at room temperature between 2 and 30 ° C, protected from light.
- Use immediately once opened and discard the excess product.

CONDITION OF SALE

Sale with Veterinary Medical prescription.

PRESENTATION

Case with 3 vials of 5 mL ampoule and Case with 1 vial of 5 mL

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

- Chile: Reg. SAG N° 530

ULTRAFIL® PLUS OTIC SUSPENSION

SUSPENSIÓN ÓTICA

ANTIINFLAMATORIO - ACARICIDA - ANTIMICÓTICO - ANTIBACTERIANO

Technical Specification

SPECIES

Dogs and cats.

ULTRAFIL® PLUS, otic suspension, is indicated in the treatment of external otitis caused by bacteria, fungi and / or mites sensitive to the association. Its use is especially indicated in ear infections caused by bacteria such as, *Staphylococcus spp., Streptococcus spp., Pseudomona aeruginosa, Proteus spp.* and *Escherichia coli*; fungi such as, *Malassezia pachydermatis, Candida spp. and Microsporum canis,* and / or mites of the *Otodectes cynotis* species.

DOSAGE FORM

Otic Suspension

THERAPEUTIC ACTION

Anti-inflammatory - Acaricide - Antifungal - Antibacterial

COMPOSITION

Each mL of suspension contains:
Gentamicin Sulfate ..... 5.09 mg
(Equivalent to 3.0 mg of Gentamicin base)
Betamethasone Dipropionate ..... 1.29 mg
(Equivalent to 1.0 mg of Betamethasone base)
Clotrimazole ..... 10 mg
Thiabendazole .....40 mg
Excipients q.s.p ..... 1 mL

PROPERTIES

ULTRAFIL® PLUS is a pharmacological association of gentamicin, clotrimazole, betamethasone and thiabendazole, which give it antibacterial, antifungal, anti-inflammatory and acaricidal properties.

Gentamicin is a bactericidal aminoglycoside antibiotic with good activity against a variety of bacteria, especially gram-negative aerobic bacilli. Gentamicin works by inhibiting normal protein synthesis in susceptible bacteria. Specifically, gentamicin is active against organisms commonly isolated in cases of otitis, such as: Staphylococcus spp., Streptococcus spp., Pseudomona aeruginosa, Proteus spp., And Escherichia coli, among others.

Clotrimazole is an imidazole antifungal agent, which is used in the treatment of infections caused by various species of pathogenic dermatophytes, fungi and Malassezia sp. Its action affects the synthesis of essential components of the fungal plasma membrane, consequently affecting their growth and division. In vitro studies have shown its activity against Candida spp., Trichophyton rubrum, Trichophyton mentagrophytes, Microsporum canis and Malassezia pachydermatis.

Betamethasone is a long-acting synthetic glucocorticoid. It has an anti-inflammatory potency 25 times more powerful than hydrocortisone, and it does not have mineralocorticoid activity. Its action decreases or prevents tissue responses to inflammatory processes in such a way that they reduce the symptoms of inflammation without affecting the basal cause.

Thiabendazole is a benzimidazole agent, with antiparasitic, antifungal and acaricidal activity. Its action occurs through the inhibition of the enzyme fumarate reductase, blocking mitochondrial function, generating an inability to obtain energy, and causing the death of the parasite. Furthermore, it is described that benzimidazoles generally bind to beta-tubulin, blocking tubulin polymerization to microtubules, consequently damaging the integrity and transport functions of parasitic cells. Studies have proven its excellent activity against the Otodectes cynotis mite in cases of otitis in dogs and cats.

INDICATIONS

- ULTRAFIL® PLUS is contraindicated in patients with hypersensitivity to any of its components.
- Do not use in patients with tympanic perforation.
- Do not use in conjunction with drugs that can induce ototoxicity.
- Do not administer during pregnancy and lactation.
- Do not administer in breeding animals.

ROUTE OD ADMINISTRATION AND DOSAGE

Topical administration by instillation into the external ear canal.

How to use and dosage:

- Shake before using.
- Clean and dry the external ear canal before applying the product.
- Remove all foreign material such as exudate, cell debris, etc.
- Trim excess hair from the area to be treated.
- Verify the integrity of the tympanic membrane.
- Instill 3 to 5 drops of Ultrafil® Plus into the ear canal to be treated, once a day, for 7 consecutive days.

CONTRAINDICATIONS

- ULTRAFIL® PLUS is contraindicated in patients with hypersensitivity to any of its components.
- Do not use in patients with tympanic perforation.
- Do not use in conjunction with drugs that can induce ototoxicity.
- Do not administer during pregnancy and lactation.
- Do not administer in breeding animals.

PRECAUTIONS

- Before starting treatment, identification of the etiologic agent (s) should be carried out, either by smear or culture. Furthermore, the antibiotic susceptibility of pathogenic bacteria should be evaluated before using this preparation.
- Before instilling any medication into the ear, examine the external ear canal to ensure there is no rupture of the tympanic membrane, in order to avoid contamination of the middle ear, as well as vestibular and cochlear injuries.
- The Dachshund breed may be particularly sensitive to treatment with Thiabendazole.

SPECIAL PRECAUTIONS FOR USE

- Before starting treatment, identification of the etiologic agent (s) should be carried out, either by smear or culture. Furthermore, the antibiotic susceptibility of pathogenic bacteria should be evaluated before using this preparation.
- Before instilling any medication into the ear, examine the external ear canal to ensure there is no rupture of the tympanic membrane, in order to avoid contamination of the middle ear, as well as vestibular and cochlear injuries.
- The Dachshund breed may be particularly sensitive to treatment with Thiabendazole.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Wash hands after administering the product.
- If the product comes into contact with the eyes, wash with plenty of water
- In case of accidental ingestion do not induce vomiting. Get immediate medical help.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

- Using ULTRAFIL® PLUS for more than 7 days may delay wound healing.
- The use of ULTRAFIL® PLUS may be associated with deafness or partial hearing loss in a small number of sensitive (geriatric) dogs. Hearing deficit is usually temporary. If vestibular or hearing dysfunctions are detected during the course of treatment, therapy should be discontinued and a flushing of the external ear canal should be performed.

OBSERVATIONS

External use only.

SECURITY

Clinical studies and safety studies of otic suspensions formulated with these compounds provide a wide margin of safety according to the recommended dose level. In cases of massive applications, patients may present symptoms associated with overdoses of each of its components, such as:

Gentamicin: When absorbed systemically it has the potential to cause nephrotoxicosis, neurotoxicosis, and ototoxicosis.
Betamethasone: Its brief administration, even in massive doses, does not usually cause harmful effects. Its chronic use can cause symptoms of hyperadrenocorticism such as polydipsia, polyuria, weight gain, sodium retention, potassium loss, etc.
Clotrimazole: It is unlikely that there will be signs of an overdose caused by clotrimazole since it is poorly absorbed after its dermal application.
Thiabendazole: A modest overdose is unlikely to cause significant problems. In dogs it has been seen that high and chronic doses could cause signs such as vomiting, diarrhea, alopecia and lethargy.

Precautions for disposing of unused product or waste material:
Do not remove empty containers or product residues on the ground or water courses. Discard the remains of unused product in its original container. Dispose of this product with caution with household waste.

CONSERVATION

- Keep at a temperature between 2 ° C and 30 ° C, away from light.
- Once opened for the first time, the duration of the product is 3 months.

CONDITION OF SALE

Sale with Veterinary Medical prescription.
Peru: Free Sale

PRESENTATION

20 mL bottle

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG Nº 2210-B
Perú: Reg. SENASA F.83.47.I.0090

COUNTRIES WHERE IT IS MARKETED

Imported and distributed in Peru by:
Representations Durand SAC. Av. Manuel Olguin
No. 501 Office No. 604 Santiago de Surco Lima.

URSOVET® ORAL SUSPENSION

SUSPENSIÓN ORAL.

COLERÉTICO, CITOPROTECTOR E INMUNOMODULADOR.

Technical Specification

SPECIES

Dogs and cats.

Ursovet® oral suspension is indicated for the treatment of inflammatory hepatobiliary diseases that occur with hepatobiliary cholestasis, in dogs and cats. Due to its choleretic, cytoprotective and immunomodulatory properties, its use is useful in cases of chronic hepatitis, cholangitis, cholangiohepatitis and cases of cholestasis that do not present with biliary obstruction.

DESCRIPTION

Choleretic, cytoprotective and immunomodulatory.

DOSAGE FORM

Oral suspension

THERAPEUTIC ACTION

Choleretic, cytoprotective and immunomodulatory.

COMPOSITION

Each 1 mL of oral suspension contains:
Ursodeoxycholic acid ..... 50 mg
Excipients c.s.p. ..... 1 mL

INDICATIONS

- Do not use in patients with biliary obstruction, fistula, pancreatitis or other complication associated with cholelithiasis.
- Do not use in patients with known hypersensitivity to bile acids.
- Do not use in rabbits or other colonic fermenting species. In these species, Ursodeoxycholic acid is converted to lithocholic acid (toxic).
- Special caution should be exercised in patients with chronic liver disease, as they may have greater difficulty in metabolizing bile acids.
- Do not use in pregnant females or during the lactation period.

ROUTE OD ADMINISTRATION AND DOSAGE

Administration way: Oral.

Dose of the active principle in dogs and cats: 10 to 15 mg / Kg, every 24 hours or divided every 12 hours.

Product dosage:

- **Dogs:** 2 to 3 mL for every 10 kg of body weight, every 24 hours or divided every 12 hours.
- **Cats:** 0.2 to 0.3 mL for each Kg of body weight, every 24 hours or divided every 12 hours.

It is recommended that treatment be administered for 3 to 4 months after the patient has presented improvement in the biochemical markers for liver disease.

DRUG INTERACTIONS

Antacids containing Aluminum or Cholestyramine resin can bind Ursodeoxycholic Acid reducing its effectiveness.

CONTRAINDICATIONS

- Do not use in patients with biliary obstruction, fistula, pancreatitis or other complication associated with cholelithiasis.
- Do not use in patients with known hypersensitivity to bile acids.
- Do not use in rabbits or other colonic fermenting species. In these species, Ursodeoxycholic acid is converted to lithocholic acid (toxic).
- Special caution should be exercised in patients with chronic liver disease, as they may have greater difficulty in metabolizing bile acids.
- Do not use in pregnant females or during the lactation period.

PRECAUTIONS

Wash your hands after administering the product.

SPECIAL PRECAUTIONS FOR USE

Wash your hands after administering the product.

WARNINGS

Mantener alejado del alcance de los niños.

ADVERSE EFFECTS

- In rare cases, there may be episodes of vomiting and / or diarrhea. In this case, discontinue use and consult your Veterinarian.
- Faced with overdose, it is recommended to perform a gastric emptying, or the administration of activated carbon or an antacid containing aluminum.

OBSERVATIONS

Shake before using.

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of water. Do not reuse the container. Dispose of the waste of this product where possible, in suitable places where special waste is disposed of.

CONSERVATION

Store at room temperature, between 15 ° and 30 ° C.
Once opened, use the product within 3 months. Discard the unused product after that period of time.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

60 mL bottle

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 2034
Costa Rica: Reg. N° MAG CL 4-48-12-6133
Uruguay: Reg. MGAP: 2018A00569
Perú: Registro SENASA F.G7.02.I.0001

COUNTRIES WHERE IT IS MARKETED

Imported in Uruguay by VIVAFIL S.A.
RIO NEGRO 1107 Montevideo - Uruguay,
TEL 29001112
grupotecnovet@gmail.com
Technical Director: DMTV Diego Cuadrado.

Imported and Distributed in Peru by Representaciones Durand SAC.
Av. Manuel Olguín No. 501 Office No. 604 Santiago de Surco Lima.

Imported and distributed in Bolivia by:
ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.

VERMIQUANTREL® ORAL TABLET

COMPRIMIDO ORAL.

ANTIPARASITARIO INTERNO TENICIDA.

Technical Specification

SPECIES

Dogs and cats

It acts on *Dipylidium caninum*, *Taenia ovis*, *Taenia multiceps*, *Taenia pisiformis, Taenia taeniformis, Taenia hydatigena, Taenia serialis and Echinococcus granulosus.*
Maximum efficiency, large safety margin.

DOSAGE FORM

Oral tablet

THERAPEUTIC ACTION

Internal antiparasitic tenicide.

COMPOSITION

Each tablet contains:
Praziquantel ..... 50 mg
Excipients q.s.p ..... 1 tablet

INDICATIONS

- Do not administer to dogs under 4 weeks of age.
- Do not administer to cats under 6 weeks of age.
- Do not administer to pregnant or lactating females.
- Do not administer to animals with hypersensitivity to Praziquantel.

ROUTE OD ADMINISTRATION AND DOSAGE

Route of Administration: Oral route

Dose of the active principle:
5 mg of Praziquantel per Kg of weight.

Product dosage:
1 tablet for every 10 kilos of weight

CONTRAINDICATIONS

- Do not administer to dogs under 4 weeks of age.
- Do not administer to cats under 6 weeks of age.
- Do not administer to pregnant or lactating females.
- Do not administer to animals with hypersensitivity to Praziquantel.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Wash your hands with soap and plenty of water after handling the product.
- In the case of contact with the eyes, it is recommended to wash with plenty of water.
- In case of accidental ingestion, do not induce vomiting. Get medical help.

WARNINGS

Mantener fuera del alcance de los niños

ADVERSE EFFECTS

In dogs, oral administration of Praziquantel can cause anorexia, vomiting, lethargy or diarrhea, but the incidence of these signs is less than 5%, while in cats adverse effects are quite rare (<2%), being reported salivation and diarrhea in these cases.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard any unused product remains in its original container. Do not throw the empty container or with product remains, in rivers, lakes or streams of natural water. Dispose of this product waste carefully with household waste.

CONSERVATION

Store in a cool and dry place, at room temperature between 15° and 30° C.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

Box with 1 tablet

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Reg. S.A.G. N°: 396

VIDATOL® INJECTABLE SOLUTION.

SOLUCIÓN INYECTABLE.

VITAMINAS DEL COMPLEJO B CON HIERRO, METIONINA Y COLINA

Technical Specification

SPECIES

Horses, cattle, pigs, sheep and goats.

Coadjuvant in the treatment of diseases that cause alterations of the nervous system, such as neuralgia, polyneuritis, neuritis and muscular dystrophy; circulatory skin disorders, such as macrocytic anemia, liver protector in convalescent animals or those with poor nutrition. It is also recommended as an appetite stimulant, in animals in training, growth, lactation or pregnancy.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Vitamins of the B complex with Iron, Methionine and Choline

COMPOSITION

Each 100 mL contains:
Thiamine Hydrochloride ..... 1.0 g
Riboflavin 5 Sodium Phosphate ..... 0.5 g
Nicotinamide ..... 5.0 g
Pyridoxine Hydrochloride ..... 0.3 g
Cyanocobalamin ..... 1.5 mg
D, L Methionine ..... 1.0 g
Choline Chloride ..... 1.0 g
Iron III Citrate hydrate ..... 4.0 g
Excipients q.s.p ..... 100 mL

INDICATIONS

Do not use in animals with a history of parenteral hypersensitivity to Thiamine (Vitamin B1), due to the possible risk of anaphylactic shock.

ROUTE OD ADMINISTRATION AND DOSAGE

The following dosages are recommended:

- **Adult cattle - adult horses:** 5 to 10 m / day, intramuscularly.
- **Sheep - pigs - goats:** 1 to 2 mL / day, intramuscularly.
- **Small cattle - small horses:** 1 to 2 mL / day intramuscularly.

Administer once a week for 1 month
In cases of horses in training or competition it is recommended to use once a week for 1 to 2 months.
In calves with neuritis symptoms or for toning it is recommended to administer once every 48 hours for four times.

CONTRAINDICATIONS

Do not use in animals with a history of parenteral hypersensitivity to Thiamine (Vitamin B1), due to the possible risk of anaphylactic shock.

WARNINGS

Mantener fuera del alcance de los niños.

GUARD PERIOD

Meat: 0 days.

Milk: 0 days.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Discard the empty container or with product remains together with household waste.
Do not reuse the container.

CONSERVATION

Keep at room temperature between 15 and 30 ° C, protected from light. Once opened, use the product within 5 weeks.
Discard the unused product after that period of time.

CONDITION OF SALE

Sale with veterinary prescription.

PRESENTATION

10 mL and 100 mL vial

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 1367
Rep. Dominicana: Reg. Nº 5604
El Salvador: VE2006053415

XILA-10® - INJECTABLE SOLUTION

SOLUCIÓN INYECTABLE.

SEDANTE-ANALGÉSICO.

Technical Specification

SPECIES

Horses and cattle.

Xila-10, solution for injection, produces a state of sedation accompanied by general muscle relaxation and a reduction in painful sensations. Due to its pharmacological properties it is indicated in:

- Handling and transporting excessively nervous or aggressive animals.
- Diagnostic procedures.
- Dental procedures
- Short-term minor surgeries, such as: cleaning and suturing of wounds, removal of dermal neoplasms, etc.
- Orthopedic procedures such as trimming and hardware.
- Pre-anesthesia in major or prolonged surgeries, with general or local anesthetics.

DOSAGE FORM

Injectable solution.

THERAPEUTIC ACTION

Sedative - Analgesic

COMPOSITION

Each 1 mL of solution contains:
Xylazine Hydrochloride ..... 117 mg
(equivalent to 100 mg of Xylazine base)
Excipients q.s.p ..... 1 mL

INDICATIONS

- Do not use concomitantly with adrenoreceptor stimulants (Adrenaline, Noradrenaline, Dopamine).
- Do not administer to animals with ventricular arrhythmias.
- Do not use in animals with intestinal obstruction or intestinal impaction.
- Do not administer in pregnant females in the last third of gestation, except during delivery.
- Do not administer in lactating females.

MODE OF APPLICATION

Intravenous administration should be done slowly.
After the injection of Xila-10®, the animal should be kept at rest until the desired effect has been achieved.

DRUG INTERACTIONS

Xylazine should not be used in joint therapy with neuroleptics or tranquilizers. The use of Xylazine and barbiturates causes additive depressant effects; the use of barbiturates to induce anesthesia should be at a reduced dose level and administered slowly when injected intravenously.
Xylazine should not be used concomitantly with adrenoreceptor stimulants (Adrenaline, Noradrenaline, Dopamine).

CONTRAINDICATIONS

- Do not use concomitantly with adrenoreceptor stimulants (Adrenaline, Noradrenaline, Dopamine).
- Do not administer to animals with ventricular arrhythmias.
- Do not use in animals with intestinal obstruction or intestinal impaction.
- Do not administer in pregnant females in the last third of gestation, except during delivery.
- Do not administer in lactating females.

PRECAUTIONS

Special warnings and precautions for use:

- To achieve the desired effect, during the administration of the drug the animal must be calm and fasting for at least 6 hours.
- Use with caution in animals with lung conditions, liver or kidney dysfunction, severe heart disease, shock, or that are under stress conditions, such as extreme heat or cold, at altitude, fatigued.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR USE

Special warnings and precautions for use:

- To achieve the desired effect, during the administration of the drug the animal must be calm and fasting for at least 6 hours.
- Use with caution in animals with lung conditions, liver or kidney dysfunction, severe heart disease, shock, or that are under stress conditions, such as extreme heat or cold, at altitude, fatigued.
- Keep out of the reach of children.

SPECIAL PRECAUTIONS FOR THE OPERATOR

- Wash your hands after administering the product.
- While handling the product, do not eat, drink liquids, or smoke.
- Avoid contact with skin and eyes.
- In the case of contact with the eyes, it is recommended to wash with plenty of water for 15 minutes.
- In case of accidental ingestion, call a doctor or a poison information center. Do not induce vomiting.
- In case of accidental self-injection, consult a doctor immediately.

WARNINGS

Mantener fuera del alcance de los niños.

ADVERSE EFFECTS

- In horses, at therapeutic doses it can occasionally cause mild muscle tremor, bradycardia with partial AV block, and a decrease in respiratory rate.
- In cattle, profuse salivation and ruminal atony may occur.

GUARD PERIOD

Meat: 2 days. Do not administer to horses whose meat is intended for human consumption.

Milk: 2 days.

OBSERVATIONS

Special precautions for disposal of unused product or waste material:
Dispose of the waste of this product with care together with household waste. Do not dispose of containers with product remains on the ground or water courses. For expired or unused products, contact the manufacturing laboratory.

CONSERVATION

Store between 15 ° and 30 ° C, protected from light.

CONDITION OF SALE

Sale only with veterinary prescription.

PRESENTATION

50 mL vial

PREPARED BY

Laboratorio Drag Pharma Chile Invetec S.A.

RECORDS

Chile: Reg. SAG N° 1837
Rep. Dominicana: Reg. N° 9102

AVAILABLE FOR SALE

Imported and distributed in Bolivia by:
ZOOFARMA
TEL: +(591)222-3357
Street Díaz Romero 1339, La Paz.