COVID-19 ALERT CONCERNING DURVET IVERMECTIN PRODUCTS: WARNING!

A number of Durvet products including Duramectin Equine Dewormer, Ivermectin Equine Dewormer, Ivermectin Sheep Drench, Ivermectin Pour-On, Ivermectin Injection and Ivermectin Plus Injection contain the anti-parasite active ingredient Ivermectin. Despite media reports that Ivermectin could potentially be used to treat people with COVID-19, these products are not safe or approved for human use, which could cause severe personal injury or death.







Keep out of reach of children.

For Oral Use In Horses Only

DURAMECTIN (ivermectin) **PASTE** 1.87%

Anthelmintic & Boticide

Net Wt. 0.21 oz (6.08 g)

Removes worms and bots with a single dose. Contents will treat up to 1250 lb body weight. ANADA# 200-326, Approved by FDA

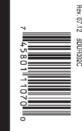
Removes Worms & Bots





Anthelmintic & Boticide

Removes Worms & Bots



INDICATIONS: Consult your veterinarian for assistance in the diagnosis, treatment and control of parasitism. DuraMectin™ (ivermectin) Paste provides effective treatment and control of the following parasites in horses. Large Strongyles (adults)- Strongylus horse's head for a few seconds after dosing. vulgaris (also early forms in blood vessels), S. edentatus (also tissue stages), S. equinus, Triodontophorus spp. including T. brevicauda and T. serratus and Craterostomum acuticaudatum; Small Strongyles (adults, including those resistant to some benzimidazole class compounds)- Coronocyclus spp. including C. coronatus, C. labiatus and C. labratus, Cyathostomum spp. including C. catinatum and C. pateratum; Cylicocyclus spp. including C insigne, C leptostomum, C nassatus, and C brevicapsulatus; Cylicodontophorus spp.; Cylicostephanus spp. including C. calicatus, C. goldi, C. longibursatus, and C. minutus and Petrovinema poculatum, Small Strongyles-Fourth-stage larvae; Pinworms (adults and fourth-stage larvae) - Oxyuns equi; Ascarids (adults, third-stage and fourth-stage larvae)

Dictyocaulus amfieldi; Intestinal Threadworms (adults) - Strongyloides westeri; Summer Sores caused by Habronema and Draschia spp. cutaneous third-stage larvae; Dermatitis caused by neck threadworm microfilariae, Onchocerca sp. DOSAGE AND ADMINISTRATION: This syringe contains sufficient paste to treat one 1250 lb horse at the recommended dose rate of 91 mcg ivermectin per lb (200 mcg/kg) body weight. Each weight marking on the syringe plunger delivers enough paste to treat 250

Parascaris equorum; Hairworms (adults)- Trichostrongylus axei, Large-mouth Stomach Worms (adults)- Habronema muscae, Bots (oral and gastric stages)- Gasterophilus spp. including G. intestinalis and G. nasalis; Lungworms (adults and fourth-stage larvae)-

Ib body weight. 👼 (1) While holding plunger, turn the knurled ring on the plunger 1/4 turn to the left and slide it so the side nearest the barrel is at the prescribed weight marking. (2) Lock the ring in place by making a 1/4 turn to the right. (3) Make sure that the horse's mouth contains no feed. (4) Remove the cover from the tip of the syringe. (5) Insert the syringe tip into the horse's mouth at the space

between the teeth. (6) Depress the plunger as far as it will go, depositing paste on the back of the tongue. (7) Imm

PARASITE CONTROL PROGRAM: All horses should be included in a regular parasite control program with particular attention being paid to mares, foals and yearlings. Foals should be treated initially at 6 to 8 weeks of age, and routine treatment releated as appropriate. Consult your veterinarian for a control program to meet your specific needs. DuraMectin™ (ivermect) Paste effectively controls gastrointestinal nematodes and bots of horses. Regular treatment will reduce the chances of v rminous arteritis caused by Strongylus vulgaris.

PRODUCT ADVANTAGES: Broad-spectrum Control - DuraMectin™ (ivermectin) Paste kills important internal parases, including bots and the arterial stages of S. vulgaris, with a single dose. DuraMectin™ (ivermectin) Paste is a potent antiparasitic agent that is neither a benzimidazole nor an organopho

ANIMAL SAFETY: DuraMectin™ (ivermectin) Paste may be used in horses of all ages, including mares at any stage of pregnancy.

WARNING: Do not use in horses intended for human consumption.

Not for use in humans. Keep this and all drugs out of reach of children. Refrain from smoking and eating when handling. Wash hands after use. Avoid contact with eyes. The Material Safety Data Sheet (MSDS) contains more detailed occupational safety information. To report adverse reactions in users, to obtain more information or to obtain MSDS, contact Durvet at 1-800-821-55

(ivermectin) PASTE 1.87% DURAMECTIN



PRECAUTIONS: DuraMectinTM ((vermectin) Paste has been formulated specifically for use in horses only. This product should not be the result of microfilariae dying in large numbers. Symptomatic treatment may be advisable. Consult your veterinarian should any used in other animal species as severe adverse reactions, including fatalities in dogs, may result

Environmental Safety: Ivermectin and excreted ivermectin residues may adversely affect aquatic organisms. Do not contaminate ground or surface water. Dispose of the syringe in an approved landfill or by incineration.

INFORMATION FOR HORSE OWNERS: Swelling and itching reactions after treatment with DuraMectin™ (ivermectin) Paste have occurred in horses carrying heavy infections of neck threadworm (Onchocerca sp. microfilariae). These reactions were most likely such reactions occur.

Healing of summer sores involving extensive tissue changes may require other appropriate therapy in conjunction with treatment with DuraMectin™ (ivermectin) Paste. Reinfection, and measures for its prevention, should also be considered. Consult your veterinarian if the condition does not improve.

STORAGE INFORMATION: STORE BETWEEN 20°C - 25°C (68°F - 77°F). Excursions between 15°C - 30°C (59°F - 86°F) are permitted.





Keep out of reach of children.

For Oral Use In Horses Only



Anthelmintic & Boticide

Removes worms and bots with a single dose.

Net Wt. 0.21 oz (6.08 g)

Contents will treat up to 1250 lb body weight ANADA# 200-326, Approved by FDA

Apple Flavored!



Anthelmintic & Boticide

Apple Flavored!



INDICATIONS: Consult your veterinarian for assistance in the diagnosis, treatment and control of parasitism. Ivermectin Paste provides effective treatment and control of the following parasites in horses. Large Strongyles (adults)- Strongylus vulgaris (also early forms in blood vessels), S. edentatus (also tissue stages), S. equinus, Triodontophorus spp. including T. brevicauda and T. serratus and Craterostornum acuticaudatum; Small Strongyles (adults, including those resistant to some benzimidazole class compounds)- Coronocyclus spp. including C. coronatus, C. labiatus and C. labratus, Cyathostornum spp. including C. catinatum and C. pateratum: Cylicocyclus spp., including C. insigne, C. leptostomum, C. nassatus, and C. brevicapsulatus; Cylicodontophorus spp.; Cylicostephanus spp. including C. calicatus, C. goldi, C. longibursatus, and C. mirutus and Petrovinema poculatum; Small Strongyles-Fourth-stage larvae; Pinworms (adults and fourth-stage larvae) - Oxyunis equi; Ascarids (adults, third-stage and fourth-stage larvae) – Parascaris equorum; **Hairworms** (adults) – Trichostrongylus axei; **Large-mouth Stomach Worms** (adults) – Habronenia muscae, **Bots** (oral and gastric stages) – Gasterophius spp, including G. intestinalis and G. nesalis, **Lungworms** (adults and fourth-stage larvae) – Dictyocaulus armifeldi; **Intestinal Threadworms** (adults) – Storogyloides westeri, **Stummer Sores** caused by

Habronema and Draschia spp. cutaneous third-stage larvae; Dermatitis caused by neck threadworm microfilariae, Onchocerca sp. DOSAGE AND ADMINISTRATION: This syringe contains sufficient paste to treat one 1250 lb horse at the recommended dose rate of 91 mcg ivermectin per lb (200 mcg/kg) body weight. Each weight marking on the syringe plunger delivers enough paste to treat 250 lb body weight.

(1) While holding plunger, turn the knurled ring on the plunger 1/4 turn to the left and slide it so the side nearest the barrel is at the prescribed weight marking. (2) Lock the ring in place by making a 1/4 turn to the right. (3) Make sure that the horse's mouth safety information. To report adverse reactions in users, to obtain more information or to obtain MSDS, contact Durvet at contains no feed. (4) Remove the cover from the tip of the syringe. (5) Insert the syringe tip into the horse's mouth at the space

between the teeth. (6) Depress the plunger as far as it will go, depositing paste on the back of the tongue. (7) In mediately raise the horse's head for a few seconds after dosing.

PARASITE CONTROL PROGRAM: All horses should be included in a regular parasite control program with particit at attention being paid to mares, foals and yearlings. Foals should be treated initially at 6 to 8 weeks of age, and routine treated as appropriate. Consult your veterinarian for a control program to meet your specific needs. Ivermectin Paste e ectively controls gastrointestinal nematodes and bots of horses. Regular treatment will reduce the chances of vermino caused by Strongylus vulgaris.

PRODUCT ADVANTAGES: Broad-spectrum Control - Ivermectin Paste kills important internal parasites, including tots and the arterial stages of S. vulgaris, with a single dose. Ivermectin Paste is a potent antiparasitic agent that is neither enzimidazole nor an organophosphate

ANIMAL SAFETY: Ivermectin Paste may be used in horses of all ages, including mares at any stage of pregnancy. Stallions may

WARNING: Do not use in horses intended for human consumption.

Not for use in humans. Keep this and all drugs out of reach of children. Refrain from smoking and eating when handling. Wash hands after use. Avoid contact with eyes. The Material Safety Data Sheet (MSDS) contains more detailed occupational 1-800-821-5570.

OBSERVE LABEL

PRECAUTIONS: Ivermectin Paste has been formulated specifically for use in horses only. This product should not be used the result of microfilariae dying in large numbers. Symptomatic treatment may be advisable. Consult your veterinarian in other animal species as severe adverse reactions, including fatalities in dogs, may result

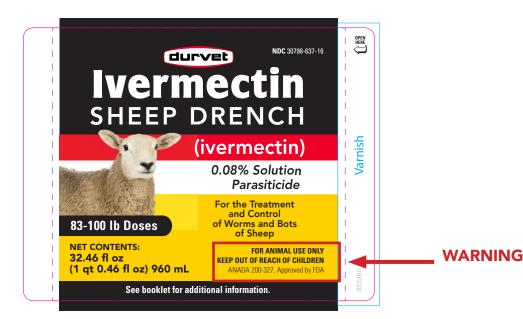
Environmental Safety: Ivermectin and excreted ivermectin residues may adversely affect aquatic organisms. Do not contaminate ground or surface water. Dispose of the syringe in an approved landfill or by incineration.

INFORMATION FOR HORSE OWNERS: Swelling and itching reactions after treatment with Ivermectin Paste have occurred in horses carrying heavy infections of neck threadworm (Onchocerca sp. microfilariae). These reactions were most likely

should any such reactions occur.

Healing of summer sores involving extensive tissue changes may require other appropriate therapy in conjunction with treatment with Ivermectin Paste. Reinfection, and measures for its prevention, should also be considered. Consult your veterinarian if the condition does not improve

STORAGE INFORMATION: STORE BETWEEN 20°C - 25°C (68°F - 77°F). Excursions between 15°C - 30°C (59°F - 86°F)



WARNING



Ivermectin Sheep Drench is stable for 24 months when stored under normal conditions. Protect from lig

I termectin Sheep Drench has been demonstrated to have a wide safety margin at the recommended do e leve and may be used in sheep of all ages. Ewes may be treated at any stage of pregnancy.

ACCEPTABILITY

Coughing may be observed in some animals during and for several minutes following drenching.

ENVIRONMENTAL SAFETY

Studies indicate that when ivermectin comes in contact with the soil, it readily and tightly binds to the id and becomes inactive over time. Free ivermectin may adversely affect fish and certain water-born orisins on which hey feed. Do not permit water unnoff from feedlost to enter lakes, streams, or ground water. Do not contaminate water by direct application or by the improper disposal of drug containers. Spills of Nerme in Sheep Drench should be contained and soaked up with absorbent towels or into loses soil, Gloves should be prevent skin exposure. All the collected materials (contaminated towels and soil), as well as all used of containers, should be placed in an impervious film bag (lastici) and disposed of by incineration or in an approved

PRECAUTIONS

Ivermectin Sheep Drench has been formulated for use in sheep ONLY. This product should not be used in other animal species as severe adverse reactions, including fatalities in dogs, may result. This product is not to be used narenterally.

Protect from light. Store at controlled room temperature between 15°-30°C (59°-86°F). Keep container tightly closed when not in use.

HUMAN SAFETY

When used as recommended in sheep, Ivermectin Sheep Drench does not pose a hazard to human health. As a routine precaution, it is advisable to wash hands after use. As with all drugs, the product should be kept out of reach of children. Contact with skin and eyes should be avoided, but protective clothing is not required.

Durvet, Inc., Blue Springs, MO 64014

FOR ORAL USE IN SHEEP ONLY

Consult your veterinarian for assistance in diagnosis, treatment, and control of parasitism.

DESCRIPTION:

Nermectin Sheep Drench is a ready-to-use, free-flowing solution of ivermectin. It is formulated to deliver the recommended dose rate of 0.2 mg ivermectin per 1 kg body weight given orally at a volume of 3.0 mL per 26 lbs body weight.

INDICATIONS:

Nermectin Sheep Drench provides treatment and control of adult and fourth-stage larvae of the following parasites: Gastrointestinal Roundworms — Haemonchus contortus, Osterdagia circumcincta, Tichostrongylus axei, Coulturiformis, Cooperia curtice, Menatodirus spathiger, N. hattus, and Desophagostonum columbianum, Lungworms — Dichocaulus filaria; and all the larval stages of Nasal Bot — Destrus ovis. It also provides treatment and control of adult forms only of the following Gastrointestinal Roundworms — Haemonchus placei, Cooperia ancophora, Strongyloides pagillosus, Desophagostomum venulosum, Trichuris ovis, and Chabertia ovina.

DIRECTIONS FOR USE:

Ivermectin Sheep Drench may be used in any standard drenching equipment or in any equipment which provides a consistent dose volume. Ivermectin Sheep Drench is administered orally at a dose of 3.0 mL [2.4 mg ivermectin) per 26 lbs body weight or 200 mcg ivermectin per kilogram of body weight.

Coughing may be observed in some animals during and for several minutes following drenching.

KEEP THIS AND ALL DRUGS OUT OF REACH OF CHILDREN.

Refrain from smoking and eating when handling. Avoid contact with eyes. Immediately wash hands and any spills on the skin with plenty of soap and water following use.

PACKAGING-

Incurrent Newmetrin Sheep Drench is available in a convenient pack size: The 32.46 fl oz (960 mL) pack contains sufficient solution to treat 83 sheep averaging 100 lbs. The packs are high-density polyethylene containers, colored opaque white, with screw caps and tamper-evident seals. They are shipped in cardboard cartons as follows: 4 x 960 mL containers nor case

Restricted Drug (California) - Use Only as Directed





ISS15XB11

4

Manufactured for: **Durvet Inc.**Blue Springs, Missouri 64014

ANADA 200-237

6



See booklet for additional information.



WARNING WARNING



Consult your veterinarian for assistance in the diagnosis, treatment and control of parasitism. INDICATIONS:

For the treatment and control of gastrointestinal roundworms (including inhibited *Ostertagia ostertagi*), lungworms, grubs, horn flies, sucking and biting lice, and sarcoptic mange mites in cattle.

See package insert for complete indications and use directions.

DOSAGE:

The dose rate is 1 mL for each 22 lb of body weight. The formulation should be applied along the topline in a narrow strip extending from the withers to the tailhead.

ADMINISTRATION:

Connect the applicator gun to the back pack as follows:

Attach the open end of the draw-off tubing to the dosing equipment. (Because of the solvents used in the formulation, only the Protector Drench Gun from Instruments Supplies Limited, or equivalent, is recommended. Other applicators may exhibit compatibility problems resulting in locking, incorrect dosage or leakage.)

Replace the shipping cap with the draw-off cap and tighten down. Attach draw-off tubing to the draw-off cap.

Gently prime the applicator gun, checking for leaks.

Follow the manufacturer's direction for adjusting the dose.

When the interval between uses of the applicator gun is expected to exceed 12 hours, disconnect the gun and draw-off tubing from the product container and empty the product from the gun and tubing back into the product container. To prevent removal of special lubricants from the Protector Drench Gun, the gun and tubing must not be washed.

durvet

(ivermectin topical solution)

for Cattle contains 5 mg ivermecting

ANIMAL SAFETY. Studies conduin the U.S.A. have demonstrated disafety margin for ivermectin. Basil or plasma levels, the topically applic formulation is expected to be at lest as well tolerated by breeding anials as is the subcutaneous formulation is as is the subcutaneous formulation is performance.

RESIDUE WARNING: Cattle must not be treated within 4 days of slaughter for human consumption. Because a withdrawal time in milk has not been established, do no use in female dairy cattle of breeding age. A withdrawal period has not been established for this product in pruminating calves. Do not use in calves to be not expenses of the visual control of the product in previous of the calves of

WARNING: KEEP OUT OF REACH OF CHILDREN NOT FOR USE IN HUMANS.

and product strout not be appear or self or others because it may be also absorbed through the skin. To milmitre accidental skin contact, the user should wear a long-sleeved shirt and rubber gloves. If accidental skin contact occurs, wash immediately with soap and water. If accidental eye exposure occurs, flush eyes immediately with water and seek medical attention.

TAKE TIME OBSERVE LA

WARNING! FLAMMABLE! KEEP AWAY FROM HEAT, SPARKS, OPEN FLAME, AND OTHER SOURCES OF IGNITION. NDC 30798-840-78



IVERMECTIN

(ivermectin topical solution)



POUR-ON for Cattle

Contains 5 mg ivermectin/mL

PARASITICIDE

Kills: Roundworms (including Brown Stomach Worm), Lungworms, Grubs, Sucking Lice, Biting Lice, Mange Mites, Horn Flies

WARNING! FLAMMABLE! KEEP AWAY FROM HEAT, SPARKS, OPEN FLAME AND OTHER SOURCES OF IGNITION. WARNING! NOT FOR USE IN HUMANS.

> Restricted Drug (California) - Use Only as Directed ANADA 200-318, Approved by FDA

NET CONTENTS: 2.5 Liter (84.5 fl oz) Contains 100-550 lb Doses IVERMECTIN (ivermectin topical solution)

durvet

POUR-ON for Cattle
Contains 5 mg ivermectin/m

The topically applied formulation of IVERMECTIN (ivermectin topical solution) Pour-On delivers internal and external parasite control in one convenient low-volume application.

PRECAUTIONS:

Store at 20°C - 25°C (68°F - 77°F), with the excursion information included on the package insert and protect from light.

Use only in well-ventilated areas or outdoors.

Close container tightly when not in

Cattle should not be treated when hair or hide is wet since reduced efficacy may be experienced.

Do not use when rain is expected to wet cattle within six hours after treatment.

This product is for application to skin surface only. Do not give orally or parenterally.

Cloudiness in the formulation may occur when IVERMECTIN (ivermectin topical solution) Pour-On is stored at temperatures below 0°C (32°P). Allowing to warm at room temperature will restore the normal appearance without affecting efficacy.

Antiparasitic activity of ivermectin will be impaired if the formulation is applied to areas of the skin with mange scabs or lesions, or with dermatoses or adherent materials, e.g., caked mud or manure.

Ivermectin has been associated with adverse reactions in sensitive dogs; therefore, IVERMECTIN (ivermectin topical solution) Pour-On is not recommended for use in species other than cattle.

Do not contaminate water by direct application or by the improper disposal of drug containers. Dispose of containers in an approved landfill or by indicated in

Product No. 1IVE013 8IVE059 Rev: 07/19



Manufactured for: Durvet, Inc. Blue Springs, MO 64014 www.durvet.com





NDC 30368-844-13

NDC 30798-844-13



1% Sterile Solution

A Parasiticide for the Treatment and Control of Internal and **External Parasites of Cattle** and Swine.

> Restricted Drug (California) -Use Only as Directed.

ANADA 200-447, Approved by FDA

Net Contents: 250 mL (8.45 fl oz)

Consult your veterinarian for assistance in the diagnosis, treatment, and control of parasitism.

INDICATIONS

Cattle: For the treatment and control of gastrointestinal roundworms (including inhibited Ostertagia ostertagi), lungworms, grubs (note insert precautions), sucking lice, and mange mites in cattle.

Swine: For the treatment and control of gastrointestinal roundworms, lungworms, lice, and mange mites in swine.

RECOMMENDED DOSE

Cattle: 1 mL per 110 lb bodyweight Swine: 1 mL per 75 lb bodyweight

See package insert for complete indications, precautions, warnings, residue information, and use directions.

Do not contaminate water by direct application or by improper disposal of drug containers. Dispose of containers in an approved landfill or by incineration.

Store at 20°C to 25°C (68°F to 77°F). Protect from light.

PLACEMENT OF LOT & EXPIRY

NDC 30798-844-13



1% Sterile Solution

A Parasiticide for the Treatment and Control of Internal and **External Parasites of Cattle** and Swine.

> Restricted Drug (California) -Use Only as Directed.

ANADA 200-447, Approved by FDA

Net Contents: 250 mL (8.45 fl oz)

WARNING - NOT FOR USE IN HUMANS. Keep this and all drugs out of the reach of children.

RESIDUE WARNING: Do not treat cattle within 35 days of slaughter. Because a withdrawal time in milk has not been established, do not use in female dairy cattle of breeding age. A withdrawal period has not been established for this product in pre-ruminating calves. Do not use in calves to be processed for yeal. Do not treat swine within 18 days of slaughter.

PRECAUTIONS

Use automatic syringe equipment only. For subcutaneous injection in cattle and swine only.

Ivermectin (ivermectin) Injection has been developed specifically for use in cattle, swine, reindeer, and American bison only. This product should not be used in other animal species as severe adverse reactions, including fatalities in dogs, may result.

Manufactured by: Bimeda-MTC Animal Health Inc. Durvet, Inc. Cambridge, ON N3C 2W4 www.bimeda.com

Manufactured for: Blue Springs, MO 64014 www.durvet.com

WARNING





Ivermectin PLUS

(ivermectin and clorsulon)

INJECTION FOR CATTLE

Treats 50-550 lb Cattle

For the treatment and control of internal parasites, including adult liver flukes, and external parasites

Approved by FDA under ANADA # 200-450



250 mL

Consult your veterinarian for assistance in the diagnosis, treatment, and control of parasitism.

INDICATIONS

Ivermectin Plus (ivermectin and clorsulon) is indicated for the treatment and control of gastrointestinal roundworms (including inhibited Ostertagia ostertagi larvae), lungworms, adult liver flukes, grubs (note insert precautions), sucking lice, and mange mites (cattle scab [note insert indications]). See package insert for complete indications and use

The dosage level of clorsulon supplied by Ivermectin Plus is effective only against adult liver flukes (Fasciola hepatica).

RECOMMENDED DOSE

Ivermectin Plus should be given only by subcutaneous injection at a dose volume of 1 mL per 110 lb (50 kg) body weight. This volume will deliver 10 mg ivermectin and 100 mg clorsulon.

This bottle is designed for use with automatic syringe equipment only. It contains enough solution to treat fifty 550 lb cattle.

For example:

Body Weight (lb)	Dose (mL)	Doses per Pack
220	2	125
330	3	83
440	4	62
550	5	50
660	6	41
770	7	35
880	8	31
990	9	27
1100	10	25

For body weights over 1100 lbs., administer at 1 mL per 110 lbs.

Divide doses greater than 10 mL between two injection sites to reduce occasional transitory discomfort or site reaction.

> Coating and Ink Free area

Manufactured for: Durvet, Inc., Blue Springs, MO 64014 www.durvet.com

durvet

Ivermectin **PLUS**

(ivermectin and clorsulon)

INJECTION FOR CATTLE

Treats 50-550 lb Cattle

For the treatment and control of internal parasites, including adult liver flukes, and external parasites

Approved by FDA under ANADA # 200-450



250 mL

Ivermectin Plus (ivermectin and clorsulon) combines ivermectin for internal and external parasite control and clorsulon, which effectively controls adult liver

1% w/v ivermectin and 10% w/v clorsulon in a

WARNING: NOT FOR USE IN HUMANS. Keep this and all drugs out of the reach of children.

KESIDUE WAKNING: Do not treat cattle Within 21 days of slaughter. Because a withdrawal period in milk has not been established, do not use in female dairy cattle of breeding age. A withdrawal period has not been established for this product in pre-ruminating calves. Do not use in calves to be processed for yeal.

PRECAUTIONS

Use automatic syringe equipment only. For subcutaneous injection in cattle only. This product is not for intravenous or intramuscular use.

Protect product from light.

Restricted Drug (California) - Use Only as Directed. Ivermectin Plus (ivermectin and clorsulon) Injection for Cattle has been developed specifically for use in cattle only. This product should not be used in other animal species as severe adverse reactions, including fatalities in dogs, may result. Do not contaminate water by direct application or by improper disposal of drug containers. Dispose of containers in an approved landfill or by incineration.

ANIMAL SAFETY

In breeding animals (bulls and cows), ivermectin and clorsulon used at the recommended level had no effect on breeding performance.

Store at 20°C to 25°C (68°F to 77°F).

Use within 28 days of first puncture and puncture a maximum of 14 times with a dosage delivery device.

Product No. 1IVE051 8IVE061 Rev 07/19

