100 mg of tulathromycin/mL

For subcutaneous injection in beef and non-lactating dairy cattle and intramuscular injection in swine only. Not for use in female dairy cattle 20 months of age or older or in calves to be processed for veal.

Indications

Beef and Non-lactating Dairy Cattle

BRD – DRAXXIN Injectable Solution is indicated for the treatment of bovine respiratory disease (BRD) associated with Mannheimia haemolytica, Pasteurella multocida, Histophilus somni, and Mycoplasma hyopneumoniae; and for the control of respiratory disease in cattle at high risk of developing SRD associated with Mannheimia haemolytica, Pasteurella multocida, Histophilus somni, and Mycoplasma bovis.

IBK – DRAXXIN Injectable Solution is indicated for the treatment of infectious bovine keratoconjunctivitis (IBK) associated with Mycoplasma bovis.

Foot Rot – DRAXXIN Injectable Solution is indicated for the treatment of foot rot in cattle caused by Fusobacterium necrophorum.

Swine

DRAXXIN Injectable Solution is indicated for the treatment of swine respiratory disease (SRD) associated with Actinobacillus pleuropneumoniae, Pasteurella multocida, Bordetella bronchiseptica, Haemophilus parasuis, and Mycoplasma hyopneumoniae; and for the control of SRD associated with Actinobacillus pleuropneumoniae, Pasteurella multocida, and Mycoplasma hyopneumoniae in gilts where SRD has been diagnosed.

Dosage and Administration

Cattle

Inject subcutaneously as a single dose in the neck at a dosage of 2.5 mg/kg (1.1 mL/100 lb) body weight (BW). Do not inject more than 10 mL per injection site.

Swine

Inject intramuscularly as a single dose in the neck at a dosage of 2.5 mg/kg (0.25 mL/22 lb) BW. Do not inject more than 2.5 mL per injection site.

Contraindications

The use of DRAXXIN Injectable Solution in contraindicated in animals previously found to be hypersensitive to the drug.

Warnings

For Use in Animals Only

Not for Human Use

Keep Out of Reach of Children. Not for Use in Chickens or Turkeys.

Rescue Warnings

Cattle

Cattle intended for human consumption must not be slaughtered within 18 days from the last treatment. Do not use in female dairy cattle 20 months of age or older. A withdrawal period has not been established for the product in pre-ruminating calves. Do not use in calves to be processed for veal.

Swine

Swine intended for human consumption must not be slaughtered within 5 days from the last treatment.

Precautions

Cattle

The effects of DRAXXIN on bovine reproductive performance, pregnancy, and lactation have not been determined. Subcutaneous injection can cause a transient local tissue reaction that may result in trim loss of edible tissue at slaughter.

Swine

The effects of DRAXXIN on porcine reproductive performance, pregnancy, and lactation have not been determined. Intramuscular injection can cause a transient local tissue reaction that may result in trim loss of edible tissue at slaughter.

Adverse Reactions

Cattle

In one field study, two calves treated with DRAXXIN at 2.5 mg/kg BW exhibited transient dyspnea. Of those calves, one exhibited transient dyspnea which may have been related to pneumonia.

Swine

In one field study, one out of 40 pigs treated with DRAXXIN at 2.5 mg/kg BW exhibited mild salivation that resolved in less than four hours.

Storage Conditions

Store at or below 25°C (77°F).

How Supplied

DRAXXIN Injectable Solution is available in the following package sizes: 50 mL vial, 100 mL vial, 250 mL vial, 500 mL vial

NADA 141-244, Approved by FDA


NADA 141-244. Approved by FDA

Dosability

Animal Health

Distributed by:

Division of Pfizer Inc, NY, NY 10017

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Only one antibiotic has been shown to lower bovine respiratory disease (BRD) chronics and mortalities by more than two-thirds, and cut your BRD re-treats by half. * To see for yourself what these numbers would mean to your operation, go to www.draxxin.com/value.

Do not use in calves to be processed for veal. A pre-slaughter withdrawal time has not been determined for pre-ruminating calves. Effects on reproductive performance, pregnancy and lactation have not been determined. DRAXXIN® (tulathromycin) Injectable Solution has a pre-slaughter withdrawal time of 18 days.

*Cutoff Your Chronics and Mortalities By 70%.*