

Trinacol™ Injection

Presentation

A sterile clear yellow aqueous solution for parenteral administration.

Each ml contains Sulfadiazine BP 200 mg, Trimethoprim BP 40 mg and Chlorocresol BP 1 mg (preservative).

Uses

Trinacol is indicated in the treatment of acute, subacute and chronic conditions of bacterial origin in cattle, pigs and dogs.

The therapeutic spectrum includes both Gram-negative and Gram-positive bacteria including *Streptococci*, *Staphylococci*, *Actinobacilli*, *Actinomycae*, *Salmonella*, *Pasteurella*, *Pneumococci*, *Proteus*, *E. coli*, *Corynebacteria*, *Vibrio*, *Bordetella*, *Brucella*, *Klebsiellae* and *Haemophilae*. It is also indicated in species where there may be an existing antibiotic drug resistance. Trinacol may be administered in respiratory infections of bacterial origin including rhinitis, pneumonia, bronchitis, and in bacterial infections secondary to viral disease such as viral pneumonia or mycoplasma infection. It is also indicated in urogenital tract infections (cystitis, vaginitis, urethritis, nephritis and metritis) and alimentary tract infections (including neonatal diarrhoea and salmonellosis). Other infections include foul-in-the-foot, severe mastitis, bacterial agalactia of sows, and infections of the eye, ear and mouth.

Dosage and Administration:

Cattle and Pigs

1 ml per 16 kg body weight daily by intramuscular or slow intravenous injection.

Dogs

1 ml per 8 kg body weight daily by subcutaneous injection only. The recommended site in dogs is the loose skin at the top of the neck.

A single injection may be sufficient in uncomplicated conditions, but in severe infections treatment may be repeated up to a maximum of 5 days, or until 2 days after the symptoms resolve.

WITHDRAWAL PERIOD START

Withdrawal Period Information - PLEASE DO NOT REMOVE

Include in WP Book (yes/no)	Changed since last WP Book (yes/no)	Cattle Milk	Cattle Meat	Sheep Meat	Pig Meat	Poultry Meat	Poultry Eggs	Fish Meat	Notes
Yes	No	48 hrs	12 days		20 days	-	-	-	

WITHDRAWAL PERIOD END

Contra-indications, warnings etc

Anaphylactic shock has been observed on rare occasions following administration of potentiated sulphonamide preparations particularly by the intravenous route. For intravenous administration the product should be warmed to body temperature and injected slowly over as long a period as is reasonably practical. At the first sign of intolerance the injection should be interrupted and shock treatment initiated.

Injections should not be given by routes other than those recommended. Not to be administered intraperitoneally.

Do not administer to animals with known sulphonamide sensitivity, severe liver parenchymal damage, or blood dyscrasias.

Milk for human consumption may be taken only from cattle after 48 hours from the last treatment. Cattle intended for human consumption should not be slaughtered until 12 days after the last treatment. Pigs intended for human consumption should not be slaughtered until 20 days after the last treatment.

Adequate drinking water should be available during the therapeutic effect of the product.

Following withdrawal of the first dose, use the product within 28 days. Unused material should be disposed of in accordance with local environmental health procedures.

For Animal Treatment Only

Pharmaceutical precautions

Store at room temperature, not exceeding 25°C. Do not freeze. Crystallisation of the product at low temperatures can be reversed by gentle warming. Protect from light. Keep out of reach of children.

Legal Category

POM.

Package Quantities

100 ml.

Further Information

Nil.

Marketing authorisation number

VM 00015/4024.