FOR ANIMAL TREATMENT ONLY
KEEP OUT OF REACH OF CHILDREN

Toltrox

ORAL SUSPENSION

ACTIVE CONSTITUENT:
50 mg/mL toltrazuril

An oral suspension for the treatment and prevention of coccidiosis in cattle up to 9 months caused by *Eimeria bovis* or *Eimeria zuernii.*

1 litre

LA12260

AgriHealth
Evidence based vet medicines

TOLTROX controls coccidiosis caused by *Eimeria bovis* or *Eimeria zuernii* in young cattle (calves). Coccidiosis typically occurs in cattle less than a year old causing diarrhoea which is often blood stained. Recently weaned calves are particularly susceptible to outbreaks of clinical disease. TOLTROX is effective in one dose because it attacks all stages of the parasite in the animal.

DIRECTIONS FOR USE: By law the user must take due care, obtaining expert advice where necessary, to avoid unnecessary pain and distress when using the product other than as directed on the label.
Thoroughly shake bottle before use.

DOSAGE: Cattle: Administer 3mL of TOLTROX per 10kg of bodyweight by mouth. For the treatment of clinical disease, treat all affected and in-contact animals. For prevention of coccidiosis on farms with a known history, treat prior to the expected onset of clinical signs. For prevention of coccidiosis in recently weaned calves, treat at weaning time when meal feeding ceases.

GENERAL INSTRUCTIONS: Before using this product, obtain a veterinary diagnosis for cause of diarrhoea. Metaphylactic treatment of cattle as soon as clinical signs of disease are seen will reduce the impact of an outbreak by preventing further intestinal damage in affected animals and preventing non-affected animals from developing diarrhoea.
To obtain maximum benefit on farms with a history of coccidiosis, TOLTROX should be given approximately 1 week prior to the expected onset of clinical signs. Treatment of newly weaned calves at the time of meal withdrawal will control coccidiosis associated with weaning.

WITHHOLDING PERIOD:
It is an offence for users of this product to cause residues exceeding the relevant MRL in the Food Notice: Maximum Residue Levels for Agricultural Compounds.
Cattle producing **meat** and offal for human consumption must not be sold for slaughter during or within **56 days** of the last treatment.

WARNING - READ LABEL BEFORE USE

Handling precautions: Do not handle until all safety precautions have been read and understood.

Wear gloves when handling. Store locked up.

Suspected of damaging fertility of the unborn child. May cause damage to organs through prolonged or repeated exposure.

First Aid: IF ON SKIN: Wash with plenty of soap and water. If exposed or concerned or if you feel unwell, get medical advice/attention.

For advice call a doctor or the National Poisons Centre 0800 POISON (0800 764 766)

Disposal: Preferably dispose of product by use. Otherwise, dispose of product and packaging at an approved landfill or other approved facility.

Storage Instructions: Store below 25°C

NZ veterinary medicine registered pursuant to the ACVM Act 1997, No. A11401. See www.foodsafety.govt.nz for registration conditions

Registered to Chanelle Pharmaceuticals Manufacturing Ltd, Ireland.

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Batch:

Expiry:

LA12261

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