FOR ANIMAL TREATMENT ONLY
KEEP OUT OF REACH OF CHILDREN

Toltrox

ORAL SUSPENSION

ACTIVE INGREDIENT:

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

LB12260

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 3 mL of TOLTROX per 10 kg 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
Causes serious eye irritation.

Suspected of damaging fertility or the unborn child. May cause damage to organs through prolonged or repeated exposure. May cause long lasting harmful effects to aquatic life. Hazardous to soil organisms.

Handling Precautions: Read carefully and follow all instructions. Obtain special instructions before use. Do not handle until all safety precautions have been read and understood.

Do not breathe mist, vapours or spray. Wear gloves when handling and wash hands thoroughly after handling.

First Aid: IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. If eye irritation persists: get medical advice/attention. 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).

Environmental protection: Avoid release to the environment.

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

Storage Instructions: Store locked up below 25°C.

Registered pursuant to the ACVM Act 1997, No. A011401. See www.foodsafety.govt.nz for registration conditions.

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New Zealand Distributor:
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LC12261

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