

FOR ANIMAL TREATMENT ONLY
RESTRICTED VETERINARY MEDICINE

TYLOGUARD®

For the treatment of tylosin-susceptible infections in cattle, pigs, sheep and goats.

ACTIVE CONSTITUENTS

Contains 200 mg of tylosin base per ml.

INDICATIONS

- Cattle: Acute mastitis, respiratory infections (including pneumoenteritis), foot rot, calf diphtheria, metritis.
- Sheep and goats: Treatment of the early stages of peracute and acute contagious agalactia caused by *Mycoplasma agalactiae*, Caprine Pleuropneumonia caused by *Mycoplasma mycoides*.
- Pigs: Swine dysentery and Enteritis caused by *Campylobacter Coli*, Swine Erysipelas, Pneumonia, Mycoplasmal arthritis.

DOSAGE AND ADMINISTRATION

Administer by intramuscular injection in cattle, pigs, sheep and goats. In food producing animals, intramuscular injection must be in the anterior half of the neck. TYLOGUARD® may be administered by **slow** intravenous injection in cattle only. Rapid I/V injection may cause ataxia, dyspnoea and increase salivation. Use a clean, dry needle. For repeat doses, alternate the injection site.

- Cattle: 5 to 10 mg/kg bodyweight (2.5 ml – 5 ml per 100 kg) by the intramuscular or slow intravenous route, daily for no more than 5 days. If the I/V route is used, follow with an I/M dose 12 hours later to maintain therapeutic levels. If there is no response to treatment after 5 days, the diagnosis and treatment should be re-assessed.
- Pigs: 5 to 10 mg/kg bodyweight (2.5 ml – 5 ml per 100 kg) by intramuscular injection in the anterior half of the neck, daily for no more than 3 days. If there is no response to treatment after 3 days, the diagnosis and treatment should be re-assessed.
- Sheep and goats: 10 mg/kg bodyweight (2.5 ml per 50 kg) by intramuscular injection in the anterior half of the neck, daily for no more than 5 days. If there is no response to treatment after 5 days, the diagnosis and treatment should be re-assessed.

WITHHOLDING PERIODS

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.

- MILK: Milk intended for sale for human consumption must be discarded during treatment and for not less than 6 milkings or approximately 72 hours following the last treatment. Sheep and goat milk intended for sale for human consumption must be discarded during treatment and for not less than 35 days following the last treatment.
- MEAT: Animals producing meat and offal for human consumption must not be sold for slaughter either during treatment or within 21 days of the last treatment.

WARNINGS AND CONTRAINDICATIONS

The administration of antibiotics may result in the overgrowth of non-susceptible organisms. If new infections occur during treatment with this product, appropriate measures should be taken. Do not mix TYLOGUARD® with other parenteral products as this may cause precipitation of the active ingredient. There are no known contraindications to the use of TYLOGUARD®. As with any injectable product, some animals may have pain at the injection site or may become depressed following treatment. On rare occasions, pigs show oedema and protrusion of the rectal mucosa accompanied by erythema and pruritis of the skin. Discontinuation of the treatment leads to uneventful recovery.

Warning: Harmful – Keep out of reach of children. May cause eye irritation. Avoid contact with the eyes. May cause sensitisation on prolonged skin contact. When handling, wear chemical-resistant gloves. Very toxic to aquatic organisms. Avoid contamination of any water supply with product or empty container. Harmful to the soil environment. Dispose of this product only by using according to the label, or at an approved landfill or other approved facility. Dispose of the container at an approved landfill or other approved facility. Do not use for any other purpose. Do not handle until all safety precautions have been read and understood. Contains tylosin and benzyl alcohol.



EMERGENCY PHONE NUMBERS

Call National Poisons Centre on: 0800 764 766.

FIRST AID

If swallowed: DO NOT INDUCE VOMITING. If available, administer 8 heaped teaspoons of activated charcoal with 2-3 glasses of water. Lean patient forward or place on left side (head down, if possible) to maintain open airway and prevent aspiration. Refer for medical attention without delay. Eye contact: Immediately hold eyes open and wash continuously for at least 15 minutes with fresh running water. Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper and lower lids. Transport to a hospital without delay. Removal of contact lenses after an eye injury should only be undertaken by skilled personnel. Skin contact: Immediately flush body and clothes with large quantities of water. Quickly remove all contaminated clothing and footwear. Wash affected areas with water (and soap if available) for at least 15 minutes. Get medical attention if irritation develops. Inhaled: Remove to fresh air. Lay patient down. Keep warm and rested. If breathing is shallow or has stopped, ensure a clear airway and apply resuscitation. Perform CPR if necessary. Transport to a doctor or hospital. Advice to doctor: Treat symptomatically. Treat as for an actual or potential allergen.

DISPOSAL

Dispose of this product only by using according to the label, or at an approved landfill or other approved facility. Dispose of the container at an approved landfill or other approved facility. Do not use for any other purpose.

STORAGE

Store below 25°C. Do not refrigerate.

Registered pursuant to the ACVM Act 1997, No. A9713. See www.foodsafety.govt.nz for registration conditions.

Registered to and manufactured in New Zealand by:

VIRBAC NEW ZEALAND LIMITED

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Ask your **Virbac Area Sales Manager** or **veterinarian** for more information

Shaping the future
of animal health

