

PENETHAMATE HYDRIODIDE

Penethaone

Management

Manage

Contraindications:

DO NOT administer by intravenous (IV) injection. DO NOT use in animals that are

NOT use in animals rad are known to be hypersensitive to penicillins or cephalosporins. DO NOT administer to animals with renal disease including anuria or oliguria.

DO NOT administer to lagomorphs or small rodents.

Adverse Reactions:

In very rare cases, anaphylactic shock may occur.

STORAGE:

Store below 25°C. Keep away from direct sunlight.

Registered pursuant to the ACVM Act 1997, No A11828 See www.foodsafety.govt.nz for registration conditions.

Registered to:
Divasa-Farmavic, S.A.
Spain

RESTRICTED VETERINARY MEDICINE
KEEP OUT OF REACH OF CHILDREN
FOR ANIMAL TREATMENT ONLY

PenethaOne® Antibiotic Injection

PENETHAMATE HYDRIODIDE

For the treatment of mastitis, uterine and respiratory infections and footrot in cattle, caused by penicillin-sensitive Gram-positive bacteria.

Highly important antibiotic

1 x 10 MIU vial antibiotic 1 x 36 mL vial diluent



READ THE ENCLOSED LEAFLET BEFORE USING THIS PRODUCT

INDICATIONS:

PenethaOne is indicated for the treatment and control of infections caused by penicillin-sensitive gram-positive bacteria that cause mastitis, uterine and respiratory infections and footrot in cattle.

Distributed by: **AgriHealth NZ Ltd** www.agrihealth.co.nz Phone 0800 821 421

DIRECTIONS FOR USE:

Reconstitute the powder contained in the vial by adding all the contents (36mL) of the diluent supplied. Shake well after reconstitution.

Administer by intramuscular (IM) injection only, in the anterior half of the neck. The reconstituted aqueous suspension is stable for 1 day at 25°C (room temperature) or for 7 days when stored between 2 - 8°C (refrigerated). Use aseptic technique when withdrawing and administering each dose of PenethaOne.

Dosage: 1. Cattle: 5 MIU once daily (equivalent to 10,000 - 15,000 IU/kg bodyweight/day). Repeat dose daily for 1 - 5 days. 2. For mastitis in lactating dairy cattle: Use the above dosing or alternatively give 10 MIU on day one, followed by 5 MIU on day two.

A maximum dose volume of 20mL per injection site should be administered.

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.

Agricultural Compounds.

WITHHOLDING PERIOD: Not for use in bobby calves. MEAT (Cattle):

Animals producing meet or offal for human consumption must not be sold for slaughter either during treatment or within 7 days of the last treatment.

MILK (Cattle): Milk intended for sale for human consumption must be discarded during treatment and for not less than: 48 hours after the last treatment of 5 MIU daily dosage; or 60 hours after the last treatment of 10 MIU plus 5 MIU.





BN: EXP:

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