Tolflam

Anti-Inflammatory Injection for Cattle and Pigs™

ACTIVE CONSTITUENT:
Tolfenamic acid 40 mg/mL


POST CALVING TRAUMA
- Fast onset of pain management
- Helps cows recover faster from caving
- Early treatment may reduce the incidence of ‘Downer Cows’.

CALVES
For ALL of the following in calves:
- Diarrhoea / enteritis
- Respiratory disease / pneumonia
- Lameness / foot injuries
- De-horning / castration
- Fever.

OTHER INFLAMMATORY AND/OR PAINFUL CONDITIONS
- Eye injury
- Pink eye
- Metritis
- Joint injuries
- Transport or yard injuries
- Caesarean section.

PNEUMONIA AND RESPIRATORY DISEASES
- Only NSAID approved for pneumonia
- Long acting in respiratory disease
- Repeat treatment approved for serious infections
- Quickly reduces fever and normalises body temperature
- Second treatment approved for severe disease.

MASTITIS TREATMENT
- Approved IV treatment for E. coli infections
- Fast reduction in pain and inflammation of mastitis
- No milk discard when administered outside 12 hour WHP.

LAMENESS
- Proven to last the full 72 hours by IM injection
- Strong – fast onset of pain management
- Repeat treatment in sever or chronic disease.
Pharmacological properties and pharmacokinetic particulars:

Tolfenamic acid (N-(2-methyl-3-chlorophenyl) anthranilic acid) is a non-steroidal, anti-inflammatory drug (NSAID) belonging to the fenamate group. Tolfenamic acid exerts anti-inflammatory, analgesic and antipyretic activities. The anti-inflammatory activity of tolfenamic acid is mainly due to an inhibition of cyclo-oxygenase and thus a reduction of the synthesis of prostaglandins and thromboxanes, which are important inflammatory mediators.

INDICATIONS

For use by or under the direction of a registered veterinarian as an aid in the treatment of pneumonia and acute mastitis in cattle and metritis-mastitis-agalactia in pigs.

DIRECTIONS FOR USE

Restraints:

DO NOT inject cattle other than into muscle tissue high on the side of the neck. Injection of product into muscles other than as described is likely to result in residues in meat above the MRL.

Precautions:

Repeat treatments or higher doses could result in residues above the MRL, unless the label withholding period is extended. The prescribing veterinarian would need to advise on an extended withholding period. Safe use of Tolflam Injection during pregnancy has not been established.

Dosage and Administration:

Use the contents within six months of initial broaching. Discard any unused portion.

CATTLE:

Pneumonia: 2 mg/kg (1 mL per 20 kg bw) by intramuscular injection high in the neck (see diagram). Treatment may be repeated once only after 48 hours.

Mastitis: 4 mg/kg (1 mL per 10 kg bw) as a single intravenous injection.

PIGS:

Metritis-mastitis-agalactia: 2 mg/kg (1 mL per 20 kg bw) as a single intramuscular injection.

WITHHOLDING PERIODS

MEAT:

Cattle by intramuscular administration: DO NOT USE less than 10 days before slaughter for human consumption.

Cattle by intravenous administration: DO NOT USE less than 4 days before slaughter for human consumption.

Pigs by intramuscular administration: DO NOT USE less than 6 days before slaughter for human consumption.

MILK:

Cattle by intramuscular or intravenous administration:

Milk collected from cows within 12 hours (1 milking) following treatment MUST NOT BE USED for human consumption or processing, or fed to bobby calves.

EXPORT SLAUGHTER INTERVAL (ESI):

An ESI has not been established for this product. Note: Observing the meat withholding period may not be sufficient to mitigate potential risks to export trade. Trade advice should be sought from the registration holder on (02) 9517 1166 before using this product.

FIRST AID

If poisoning occurs, contact a doctor or Poisons Information Centre. Phone Australia 131 126.

STORAGE

Store below 30°C (room temperature) away from direct sunlight.

DISPOSAL

Dispose of empty containers by wrapping with paper and putting in garbage.

APVMA Approval No.: 83476/108367

Injection site