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Ketink 100 mg/ml solution for injection for cattle, pigs and horses

Authorised

- Ketoprofen

Product identification

Medicine name:
Ketink 100 mg/ml solution for injection for cattle, pigs and horses

Active substance:
Ketoprofen

Target species:
Cattle
Pig
Horse

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Ketoprofen

100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 4 day
  - Milk. 0 hour
- **Pig**
  - Meat and offal. 4 day
- **Cattle**
  - Meat and offal. 4 day
  - Milk. 0 hour

Intravenous use:

- **Cattle**
  - Meat and offal. 4 day
  - Milk. 0 hour
- **Horse**
  - Meat and offal. 4 day
  - Milk. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

- **Cattle**
  - Meat and offal. 4 day

- Milk. 0 hour

- **Horse**
  - Meat and offal. 4 day
  - Milk. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AE03

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Denmark

Available in:
Denmark

Package description:
cardboard box containing 1 vial of 100 ml
cardboard box containing 1 vial of 250 ml
cardboard box containing 6 vials of 100 ml
cardboard box containing 10 vials of 100 ml
cardboard box containing 12 vials of 100 ml
cardboard box containing 6 vials of 250 ml
cardboard box containing 10 vials of 250 ml
cardboard box containing 12 vials of 250 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Industrial Veterinaria S.A.

Marketing authorisation date:
27/02/2012

Manufacturing sites for batch release:
Industrial Veterinaria S.A.
aniMedica GmbH

Responsible authority:
Danish Medicines Agency

Authorisation number:
48707

Date of authorisation status change:
27/02/2012

Reference member state:
Spain

Procedure number:
ES/V/0175/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia France Germany Greece Hungary Ireland Italy Latvia Lithuania Malta Netherlands Poland Portugal Romania Slovakia Slovenia United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)
Published on: 22/12/2023
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Package Leaflet

Labelling

Combined File of all Documents