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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

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Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
cardboard box containing 12 vials of 250 ml
cardboard box containing 10 vials of 250 ml
cardboard box containing 6 vials of 250 ml
cardboard box containing 12 vials of 100 ml
cardboard box containing 10 vials of 100 ml
cardboard box containing 6 vials of 100 ml
cardboard box containing 1 vial of 250 ml
cardboard box containing 1 vial of 100 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:
22/03/2012

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

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401557.00.00

Date of authorisation status change:
23/03/2017

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)

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Documents

Labelling

Combined File of all Documents

English (PDF)
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