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

Authorised

- Ketoprofen

Product identification

Medicine name:
Ketosol-100, 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

Intravenous use:

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

Not authorized 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

Package description:
50 ml amber glass bottles (type II) closed with brombutyl rubber stopper and aluminium cap or flip-off cap with aluminium seal and polypropylene cover.
100 ml amber glass bottles (type II) closed with brombutyl rubber stopper and aluminium cap or flip-off cap with aluminium seal and polypropylene cover.

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:
Interchemie Werken De Adelaar Eesti AS

Marketing authorisation date:
27/10/2022

Manufacturing sites for batch release:
Interchemie Werken De Adelaar Eesti AS

Responsible authority:
Danish Medicines Agency

Authorisation number:
66761

Date of authorisation status change:
27/10/2022

Reference member state:
Estonia

Procedure number:
EE/V/0102/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Finland France Germany Greece Hungary Ireland Italy Latvia Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain Sweden

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 13/01/2026
Updated on: 14/03/2026
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