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KETOPROCEN 100 mg/ml SOLUTION FOR INJECTION

Authorised

- Ketoprofen

Product identification

Medicine name:
KETOPROCEN 100 mg/ml SOLUTION FOR INJECTION

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

Intravenous use:

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Cattle

- Meat and offal. 4 day
- Milk. 0 hour
- Meat and offal. 4 day
- Milk. 0 hour
- Meat and offal. 4 day
- Milk. 0 hour
- Meat and offal. 4 day
- Milk. 0 hour

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Horse

- Meat and offal. 4 day
- Milk. 0 hour
- Meat and offal. 4 day
- Milk. 0 hour
- Meat and offal. 4 day
- Milk. 0 hour
- Meat and offal. 4 day
- Milk. 0 hour

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Cattle

- Meat and offal. 4 day
- Milk. 0 hour
- Meat and offal. 4 day
- Milk. 0 hour
- Meat and offal. 4 day
- Milk. 0 hour

- Meat and offal. 4 day
- Milk. 0 hour

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Horse

- Meat and offal. 4 day
- Milk. 0 hour
- Meat and offal. 4 day
- Milk. 0 hour
- Meat and offal. 4 day
- Milk. 0 hour
- Meat and offal. 4 day
- Milk. 0 hour

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:
Lithuania

Package description:
box containing 10 vials of 50 ml
box containing 1 vial of 250 ml
box containing 10 vials of 100 ml
box containing 1 vial of 100 ml
box containing 10 vials of 250 ml
box containing 1 vial of 50 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:
Cenavisa S.L.

Marketing authorisation date:
8/11/2016

Manufacturing sites for batch release:
Cenavisa S.L.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
28/01/2026

Reference member state:
Spain

Procedure number:
ES/V/0267/001

Concerned member states:
Belgium Bulgaria Estonia Greece Hungary Italy Latvia Lithuania Poland Portugal Romania

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

Documents

Summary of Product Characteristics

Package Leaflet

Labelling

Combined File of all Documents

English (PDF)
Published on: 13/03/2026
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