

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:**

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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

-

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

-

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

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: 24/07/2025

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