

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:

Germany

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:

25/03/2022

Manufacturing sites for batch release:

Interchemie Werken De Adelaar Eesti AS

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

V7006969.00.00

Date of authorisation status change:

25/03/2022

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

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

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics