

Ketodolor 100 mg/ml solution for injection for horses, cattle, pigs

Authorised

- Ketoprofen

Product identification

Medicine name:

Ketodolor 100 mg/ml solution for injection for horses, cattle, pigs

Ketodolor 100 mg/ml šķīdums injekcijām zirgiem, liellopiem, cūkām

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

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Pig

- Meat and offal. 4 day

Intravenous use:

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Cattle

- Meat and offal. 1 day

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Horse

- Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AE03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Latvia

Package description:

The product is packed in amber type II glass vial of 50 ml fitted with red chlorobutyl stopper and aluminium cap.

The product is packed in amber type II glass vial of 100 ml, fitted with red chlorobutyl stopper and aluminium cap.

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:

Le Vet. Beheer B.V.

Marketing authorisation date:

30/05/2013

Manufacturing sites for batch release:

Produlab Pharma B.V.

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/DCP/13/0020

Date of authorisation status change:

30/05/2013

Reference member state:

Ireland

Procedure number:

IE/V/0621/001

Concerned member states:

Austria Belgium Czechia Denmark Estonia Finland France Hungary Iceland
Italy Latvia Lithuania Luxembourg Malta Poland Portugal Romania Slovakia
Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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