

# 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

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**Active substance:**

Ketoprofen

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**Target species:**

Cattle

Pig

Horse

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**Route of administration:**

Intramuscular use

Intravenous use

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## Product details

**Active substance and strength:**

Ketoprofen

100.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QM01AE03

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Denmark

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**Available in:**

Denmark

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Le Vet. Beheer B.V.

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**Marketing authorisation date:**

17/07/2014

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

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**Responsible authority:**

Danish Medicines Agency

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**Authorisation number:**

50759

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**Date of authorisation status change:**

17/07/2014

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0621/001

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**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)

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To consult adverse reactions on veterinary medicinal products please go to  
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## Documents

Summary of Product Characteristics

English (PDF)

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