

# KELAPROFEN 100 mg/ml

Authorised

- Ketoprofen

## Product identification

**Medicine name:**

KELAPROFEN 100 mg/ml

Ketaprofen 100 mg/ml, injekčný roztok pre hovädzí dobytok, kone a ošípané

**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

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Cattle**

- Meat and offal. no withdrawal period
- Meat and offal: IV: 1 day, IM: 2 days

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**Pig**

- Meat and offal. 2 day

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**Cattle**

- Milk. 0 hour

**Intravenous use:**

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**Cattle**

- Meat and offal. no withdrawal period
- Meat and offal: IV: 1 day, IM: 2 days

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**Horse**

- Meat and offal. 1 day

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**Cattle**

- Milk. 0 hour

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**Horse**

- Milk. no withdrawal period

Milk: Its use is not authorized in animals whose milk is used for human consumption.

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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:**

Slovakia

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**Package description:**

box containing 12 vials of 250 ml  
box containing 12 vials of 100 ml  
box containing 12 vials of 50 ml  
box containing 10 vials of 250 ml  
box containing 10 vials of 100 ml  
box containing 10 vials of 50 ml  
box containing 6 vials of 250 ml  
box containing 6 vials of 100 ml  
box containing 6 vials of 50 ml  
box containing 1 vial of 250 ml  
box containing 1 vial of 100 ml  
box containing 1 vial of 50 ml

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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:**

KELA Kempisch Laboratorium Kela Laboratoria

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**Marketing authorisation date:**

4/03/2015

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**Manufacturing sites for batch release:**

**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/001/MR/15-S

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**Date of authorisation status change:**

4/03/2015

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0160/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
France Germany Greece Hungary Ireland Italy Lithuania Luxembourg  
Netherlands Norway Poland Portugal Romania Slovakia Sweden  
United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

Published on: 24/07/2025

[Download](#)

Summary of Product Characteristics

Package Leaflet

Labelling

eu-PUAR-esv0160001-dcp-kelaprofen--100-mg-ml-en.pdf