File downloaded on 2026-10-30
**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000992611

# Labiprofen 150 mg/ml solution for injection

Authorised

- Ketoprofen

## Product identification

**Medicine name:**
Labiprofen 150 mg/ml solution for injection

**Active substance:**
Ketoprofen

**Target species:**
Pig
Cattle
Horse

**Route of administration:**
Intramuscular use
Intravenous use

## Product details

**Active substance and strength:**
Ketoprofen
150.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Solution for injection

---

**Withdrawal period by route of administration:**

**Intramuscular use:**

- **Pig**
  - Meat and offal. 3 day
- **Cattle**
  - Meat and offal. 2 day
  - Milk. 0 hour

**Intravenous use:**

- **Horse**
  - Meat and offal. 1 day
  - Milk. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

- **Cattle**
  - Meat and offal. 2 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:**

Austria

**Available in:**
Austria

**Package description:**
box containing 10 vials of 250 ml
box containing 10 vials of 100 ml
box containing 12 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

## 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:**
Labiana Life Sciences S.A.

**Marketing authorisation date:**
26/06/2023

**Manufacturing sites for batch release:**
Labiana Life Sciences S.A.

**Responsible authority:**
Austrian Agency For Health And Food Safety

**Authorisation number:**
841720

**Date of authorisation status change:**
26/06/2023

**Reference member state:**

Spain

**Procedure number:**
ES/V/0388/001

**Concerned member states:**
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Sweden United Kingdom (Northern Ireland)

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

## Documents

Combined File of all Documents

Package Leaflet

English (PDF)
Published on: 14/03/2026
Download

Labelling

English (PDF)
Published on: 14/03/2026

Download

Summary of Product Characteristics

English (PDF)
Published on: 14/03/2026
Download