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Labiprofen 150 mg/ml solution for injection

- Ketoprofen

Authorised

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:

- Romania

Available in:

- Romania

Package description:

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

- 24/06/2021

Manufacturing sites for batch release:

- Labiana Life Sciences S.A.

Responsible authority:

- Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

- 260096

Date of authorisation status change:

- 23/06/2026

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

Product information

Package Leaflet

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

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

Summary of Product Characteristics

English (PDF)
Published on: 14/03/2026
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Combined File of all Documents

English (PDF)

Published on: 13/03/2026
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Other languages (1)
Romanian (PDF)
Published on: 29/06/2026
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