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Ketiva 150 mg/ml solution for injection for cattle, pigs and horses

Authorised

- Ketoprofen

Product identification

Medicine name:
Ketiva 150 mg/ml solution for injection for cattle, pigs and horses

Active substance:
Ketoprofen

Target species:
Pig
Cattle
Horse

Route of administration:
Solution for injection

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:
Solution for injection:

- **Pig**
  - Meat and offal. 3 day
- **Cattle**
  - Meat and offal. 2 day
  - Milk. 0 day
- **Horse**
  - Milk. no withdrawal period

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

  - 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:
Sweden

Package description:
Amber glass vial type II, closed with a bromobutyl rubber stopper and aluminium pull off cap or aluminium/plastic flip off cap.
Amber glass vial type II, closed with a bromobutyl rubber stopper and aluminium pull off cap or aluminium/plastic flip off cap.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application – change in therapeutic indication(s) (Article 19(1)(a) of Regulation (EU) 2019/6)

Marketing authorisation holder:
Vetviva Richter GmbH

Marketing authorisation date:
19/03/2026

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
67946

Date of authorisation status change:
19/03/2026

Reference member state:
Portugal

Procedure number:
PT/V/0149/001

Concerned member states:
Austria Belgium Bulgaria Croatia Czechia Denmark Estonia Finland France Germany Greece Hungary Ireland Italy Latvia Lithuania Netherlands Norway Poland 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.

Combined File of all Documents