Rifen 100 mg/ml - Injektionslösung für Pferde, Rinder und Schweine

Authorised

- Ketoprofen

Product identification

Medicine name:
Rifen 100 mg/ml - Injektionslösung für Pferde, Rinder und Schweine

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

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 3 day
- **Pig**
  - Meat and offal. 4 day

Intravenous use:

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 1 day
- **Horse**
  - 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:
Poland

Available in:
Poland

Package description:
10 x 100 ml Amber glass vials type II, with bromobutyl rubber stopper type I and aluminium caps.
10 x 50 ml Amber glass vials type II, with bromobutyl rubber stopper type I and aluminium caps.
100 ml Amber glass vials type II, with bromobutyl rubber stopper type I and aluminium caps.
50 ml Amber glass vials type II, with bromobutyl rubber stopper type I and aluminium caps.

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:
Vetviva Richter GmbH

Marketing authorisation date:
23/11/2012

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2231

Date of authorisation status change:
23/11/2012

Reference member state:
Austria

Procedure number:
AT/V/0002/001

Concerned member states:
Belgium Czechia Denmark Finland France Germany Greece Hungary Ireland Italy Latvia Netherlands Poland Portugal Slovakia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Package Leaflet

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Summary of Product Characteristics

English (PDF)
Published on: 30/08/2024
Updated on: 13/03/2026
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Labelling

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Combined File of all Documents

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