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Comforion vet 100 mg/ml solution for injection for horse, cattle and swine

Authorised

- Ketoprofen

Product identification

Medicine name:
Comforion vet 100 mg/ml solution for injection for horse, cattle and swine

Active substance:
Ketoprofen

Target species:
Horse
Cattle
Pig

Route of administration:
Intravenous use
Intramuscular 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:
Intravenous use:

- **Horse**
  - Meat and offal. 4 day

Not authorized for use in animals producing milk for human consumption.

  - Milk. 0 hour

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

Intramuscular use:

- **Cattle**
  - Meat and offal. 4 day
  - Milk. 0 day

- **Pig**
  - Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AE03

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Norway

Available in:
Norway

Package description:
10 x 100 ml amber glass vial type II with bromobutyl rubber stopper type I and aluminium cap.
10 x 50 ml amber glass vial type II with bromobutyl rubber stopper type I and aluminium cap.
100 ml amber glass vial type II with bromobutyl rubber stopper type I and aluminium cap.
50 ml amber glass vial type II with bromobutyl rubber stopper type I and aluminium cap.

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:
Orion Corporation

Marketing authorisation date:
31/07/2003

Manufacturing sites for batch release:
Orion Corporation
Vetviva Richter GmbH

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
03 - 1776

Date of authorisation status change:
15/12/2008

Reference member state:
Finland

Procedure number:
FI/V/0101/001

Concerned member states:
Belgium Bulgaria Croatia Czechia Estonia France Hungary Latvia Lithuania Luxembourg Netherlands Norway Poland Romania Slovakia Slovenia Sweden

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 7/03/2023
Download

Package Leaflet

English (PDF)
Published on: 9/06/2023
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Combined File of all Documents