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Ketodolor 100 mg/ml solution for injection for horses, cattle, pigs

Authorised

- Ketoprofen

Product identification

Medicine name:
Ketodolor 100 mg/ml solution for injection for horses, cattle, pigs

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

Intravenous use:

- **Cattle**
  - Meat and offal. 1 day
  - Milk. 0 hour
- **Horse**
  - Meat and offal. 1 day
  - Milk. no withdrawal period

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AE03

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

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Authorisation status:
Valid

Authorised in:

Denmark

Package description:
The product is packed in amber type II glass vial of 100 ml, fitted with red chlorobutyl stopper and aluminium cap.
The product is packed in amber type II glass vial of 50 ml fitted with red chlorobutyl stopper 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:
Le Vet. Beheer B.V.

Marketing authorisation date:
17/07/2014

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Danish Medicines Agency

Authorisation number:
50759

Date of authorisation status change:
17/07/2014

Reference member state:
Ireland

Procedure number:
IE/V/0621/001

Concerned member states:
Austria Belgium Czechia Denmark Estonia Finland France Hungary Iceland Italy Latvia Lithuania Luxembourg Malta Poland Portugal 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

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