Nefotek 100 mg/ml Solution for Injection

Authorised

- Ketoprofen

Product identification

Medicine name:
Nefotek 100 mg/ml Solution for Injection
NEFOTEK 100 MG/ML SOLUTION INJECTABLE POUR BOVINS EQUINS ET PORCINS

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

Intravenous use:

- **Cattle**
  - Meat and offal. 4 day
  - Milk. 0 hour
- **Horse**
  - Meat and offal. 4 day
  - Milk. no withdrawal period

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

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

- **Horse**
  - Meat and offal. 4 day
  - Milk. no withdrawal period

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AE03

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

Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
cardboard box containing 1 vial of 250 ml
cardboard box containing 1 vial of 100 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:
Vetpharma Animal Health S.L.

Marketing authorisation date:

20/12/2011

Manufacturing sites for batch release:
Industrial Veterinaria S.A.

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/2942366 4/2011

Date of authorisation status change:
16/01/2017

Reference member state:
Spain

Procedure number:
ES/V/0176/001

Concerned member states:
Austria Belgium Czechia Denmark France Germany Hungary Ireland Italy Netherlands Poland Slovakia Slovenia United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 1/02/2024
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Package Leaflet

Labelling

Package Leaflet and Labelling

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