

Nefotek 100 mg/ml Solution for Injection

Authorised

- Ketoprofen

Product identification

Medicine name:

Nefotek 100 mg/ml Solution for Injection

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:

Poland

Package description:

cardboard box containing 1 vial of 100 ml
cardboard box containing 1 vial of 250 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:

13/04/2012

Manufacturing sites for batch release:

Industrial Veterinaria S.A.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

2177

Date of authorisation status change:

13/04/2012

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

Package Leaflet

English (PDF)

Published on: 1/02/2024

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Labelling

English (PDF)

Published on: 1/02/2024

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Summary of Product Characteristics

English (PDF)

Published on: 13/03/2026

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