

Enroxil Max 100 mg/ml solution for injection for cattle

Not authorised

- Enrofloxacin

Product identification

Medicine name:

Enroxil Max 100 mg/ml solution for injection for cattle

Active substance:

Enrofloxacin

Target species:

Cattle

Route of administration:

Intravenous use
Subcutaneous use

Product details

Active substance and strength:

Enrofloxacin
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intravenous use:**

-

Cattle

- Meat and offal. 4 day
- Milk. 72 hour

Subcutaneous use:

-

Cattle

- Meat and offal. 14 day
- Milk. 84 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Lithuania

Package description:

Amber glass Type 2 multi-dose vial of 100 ml with bromobutyl rubber stopper

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:

Krka d.d. Novo Mesto Eesti Filiaal

Marketing authorisation date:

6/10/2013

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/13/2182/001

Date of authorisation status change:

6/10/2013

Reference member state:

Ireland

Procedure number:

IE/V/0424/001

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

Documents

Combined File of all Documents

English (PDF)

Published on: 12/04/2026

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Summary of Product Characteristics