

Enroxil Max 100 mg/ml solution for injection for cattle

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:

Valid

Authorised in:

Czechia

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 tovarna zdravil d.d. Novo mesto

Marketing authorisation date:

9/10/2008

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/052/08-C

Date of authorisation status change:

7/01/2013

Reference member state:

Ireland

Procedure number:

IE/V/0424/001

Concerned member states:

Czechia Italy Latvia Lithuania Romania Slovakia

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: 12/04/2026

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Combined File of all Documents

Package Leaflet

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Labelling

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