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:
United Kingdom (Northern Ireland)

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:
1/08/2007

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 01656/4006

Date of authorisation status change:
27/06/2024

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