

Enrotril Max 100 mg/ml Solution for Injection for Cattle

Not
authorised

- Enrofloxacin

Product identification

Medicine name:

Enrotril Max 100 mg/ml Solution for Injection for Cattle

Active substance:

Enrofloxacin

Target species:

Cattle

Route of administration:

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:**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:

Czechia

Package description:

500 ml amber Type I glass vial with a grey bromobutyl rubber stopper and aluminium overseal. 500 ml vial sold in packs containing 12 x 500 ml.

500 ml amber Type I glass vial with a grey bromobutyl rubber stopper and aluminium overseal. 500 ml vial sold in packs containing 4 x 500 ml.

500 ml amber Type I glass vial with a grey bromobutyl rubber stopper and aluminium overseal. 500 ml vial sold in packs containing 1 x 500 ml.

250 ml amber Type I glass vials with a grey bromobutyl rubber stopper and aluminium overseal. 250 ml vial sold in packs containing 12 x 250 ml.

250 ml amber Type I glass vial with a grey bromobutyl rubber stopper and aluminium overseal. 250 ml vial sold in packs containing 4 x 250 ml.

250 ml amber Type I glass vial with a grey bromobutyl rubber stopper and aluminium overseal. 250 ml vial sold in packs containing 1 x 250 ml.

100 ml amber Type I glass vial with a grey bromobutyl rubber stopper and aluminium overseal. 100 ml vial sold in packs containing 12 x 100 ml.

100 ml amber Type I glass vial with a grey bromobutyl rubber stopper and aluminium overseal. 100 ml vial sold in packs containing 4 x 100 ml.

100 ml amber Type I glass vial with a grey bromobutyl rubber stopper and aluminium overseal. 100 ml vial sold in packs containing 1 x 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:

Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:

29/03/2011

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/015/11-C

Date of authorisation status change:

29/03/2011

Reference member state:

Ireland

Procedure number:

IE/V/0557/001

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 28/01/2022

[Download](#)

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.