

Enrotril Max 100 mg/ml Solution for Injection for Cattle

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:**

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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:

Romania

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:

15/10/2015

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

190143

Date of authorisation status change:

4/11/2019

Reference member state:

Ireland

Procedure number:

IE/V/0557/001

Concerned member states:

Austria Czechia Estonia Greece Latvia Lithuania Portugal Romania Slovakia
Slovenia Spain 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)

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