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

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Authorisation status:

Surrendered

Authorised in:

Spain

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 glas 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:
20/12/2010

Manufacturing sites for batch release:
Norbrook Manufacturing Limited
Norbrook Laboratories Limited

Responsible authority:
Spanish Agency For Medicines And Medical Devices

Authorisation number:
2231 ESP

Date of authorisation status change:
29/09/2023

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
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Labelling

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Package Leaflet

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