Enrotron 100 mg/ml Solution for injection for cattle, sheep, goats and pigs

Authorised

- Enrofloxacin

Product identification

Medicine name:
Enrotron 100 mg/ml Solution for injection for cattle, sheep, goats and pigs

Active substance:
Enrofloxacin

Target species:
Pig
Cattle
Goat
Sheep

Route of administration:
Intramuscular use
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:

Intramuscular use:

- **Pig**
  - Meat and offal. 13 day

Intravenous use:

- **Cattle**
  - Meat and offal. 5 day
  - Milk. 3 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 12 day
  - Milk. 4 day
- **Goat**
  - Meat and offal. 6 day
  - Milk. 4 day
- **Sheep**
  - Meat and offal. 4 day

- Milk. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Estonia

Available in:
Estonia

Package description:
Pack Size: 100 ml clear glass vial type I with siliconised, Teflon coated rubber stopper sealed with an aluminium cap.Cartons of 12 x 100 ml are available.
Pack Size: 100 ml clear glass vial type I with siliconised, Teflon coated rubber stopper sealed with an aluminium cap.Carton of 1 x 100 ml is available.
Pack Size: 100 ml clear glass vial type I with siliconised, uncoated bromobutyl rubber stopper, sealed with an aluminium cap.Carton of 1 x 100 ml is available.
Pack Size: 100 ml clear glass vial type I with siliconised, uncoated bromobutyl rubber stopper, sealed with an aluminium cap.Cartons of 12 x 100 ml are available.
Pack Size: 250 ml clear glass vial type I with siliconised, uncoated bromobutyl rubber stopper, sealed with an aluminium cap.Carton of 1 x 250 ml is available.
Pack Size: 250 ml clear glass vial type I with siliconised, uncoated bromobutyl rubber stopper, sealed with an aluminium cap.Cartons of 12 x 250 ml are available.

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:
aniMedica GmbH

Marketing authorisation date:
28/03/2012

Manufacturing sites for batch release:
aniMedica GmbH
Industrial Veterinaria S.A.

Responsible authority:
State Agency Of Medicines

Authorisation number:
1701

Date of authorisation status change:
28/03/2012

Reference member state:
Ireland

Procedure number:
IE/V/0270/003

Concerned member states:
Estonia Greece Hungary Italy Latvia Lithuania Poland Portugal Romania 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)
Published on: 16/01/2026
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Combined File of all Documents