Enrotron 100, 100 mg/ml solution for injection for cattle and pigs

Authorised

- Enrofloxacin

Product identification

Medicine name:
Enrotron 100, 100 mg/ml solution for injection for cattle and pigs

Active substance:
Enrofloxacin

Target species:
Cattle
Goat
Sheep
Pig

Route of administration:
Intravenous use
Subcutaneous use
Intramuscular 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. 5 day
  - Milk. 72 hour

Subcutaneous use:

- **Goat**
  - Meat and offal. 6 day
  - Milk. 96 hour
- **Sheep**
  - Meat and offal. 4 day
  - Milk. 72 hour
- **Cattle**
  - Meat and offal. 12 day
  - Milk. 96 hour

Intramuscular use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

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

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
(ID2) 1200 millilitre(s): unspecified outer container with 12 Vial (glass) each with 100 millilitre(s), closed with Stopfen (rubber)
(ID1) 100 millilitre(s): unspecified outer container with 1 Vial (glass) with 100 millilitre(s), closed with Stopfen (rubber)
(ID4) 3000 millilitre(s): unspecified outer container with 12 Vial (glass) each with 250 millilitre(s), closed with Stopfen (rubber)
(ID3) 250 millilitre(s): unspecified outer container with 1 Vial (glass) with 250 millilitre(s), closed with Stopfen (rubber)

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:
23/08/2012

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

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401622.00.00

Date of authorisation status change:
3/07/2017

Reference member state:
Germany

Procedure number:
DE/V/0147/003

Concerned member states:
Austria Slovenia

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

Documents

Combined File of all Documents

English (PDF)
Published on: 30/09/2025
Download

2401622-paren-20180221.rtf