Interflox-100, 100 mg/ml solution for injection for cattle, sheep, goats and pigs

Authorised

- Enrofloxacin

Product identification

Medicine name:
Interflox-100, 100 mg/ml solution for injection for cattle, sheep, goats and pigs
Interflox 100, 100mg/ml, Injekční roztok

Active substance:
Enrofloxacin

Target species:
Cattle
Sheep
Goat
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. 3 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 12 day
  - Milk. 4 day
- **Sheep**
  - Meat and offal. 4 day
  - Milk. 3 day
- **Goat**
  - Meat and offal. 6 day
  - Milk. 4 day

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:
Czechia

Package description:
100 ml amber glass bottles (type I) closed with brombutyl rubber stopper and aluminium cap in cardboard box.

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:
Interchemie Werken De Adelaar Eesti AS

Marketing authorisation date:
13/05/2019

Manufacturing sites for batch release:
Interchemie Werken De Adelaar Eesti AS

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/030/19-C

Date of authorisation status change:
13/05/2019

Reference member state:
Estonia

Procedure number:
EE/V/0103/001

Concerned member states:
Austria Bulgaria Croatia Cyprus Czechia France Greece Hungary Italy Latvia Malta Poland Portugal Romania Slovakia Slovenia Spain

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

Documents

Summary of Product Characteristics

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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.