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 mg/ml SOLUCION INYECTABLE PARA BOVINO, OVINO, CAPRINO Y PORCINO

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:
Spain

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:
25/03/2019

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

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:

3768 ESP

Date of authorisation status change:
14/02/2022

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