Baytril 100 mg/ml solution for injection

Authorised

- Enrofloxacin

Product identification

Medicine name:
Baytril 100 mg/ml solution for injection
Baytril 100 mg/ml инжекционен разтвор

Active substance:
Enrofloxacin

Target species:
Sheep
Cattle
Goat
Pig

Route of administration:
Subcutaneous use
Intravenous use
Subcutaneous 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:

Subcutaneous use:

- **Sheep**
  - Meat and offal. 4 day
  - Milk. 3 day

Intravenous use:

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

Subcutaneous use:

- **Cattle**
  - Meat and offal. 12 day
  - Milk. 4 day

Subcutaneous use:

- **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:
Bulgaria

Available in:
Bulgaria

Package description:
Brown glass (type I) vials with a chlorobutyl polytetrafluoroethylene (PTFE) stopper and with a flip-off cap with aluminium case and plastic flip-off button.
Brown glass (type I) vials with a chlorobutyl polytetrafluoroethylene (PTFE) stopper and with a flip-off cap with aluminium case and plastic flip-off button.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Elanco Animal Health GmbH

Marketing authorisation date:

21/03/2016

Manufacturing sites for batch release:
KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2637

Date of authorisation status change:
21/03/2016

Reference member state:
Hungary

Procedure number:
HU/V/0126/001

Concerned member states:
Bulgaria

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 and Labelling

This document does not exist in this language (English). You can find it in another language below.