BOFLOX 100 mg/ml

Authorised

- Marbofloxacin

Product identification

Medicine name:
BOFLOX 100 mg/ml

Active substance:
Marbofloxacin

Target species:
Cattle
Pig (sow)

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Marbofloxacin
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. no withdrawal period

Meat and offal: 6 days 2 mg (IV/SC/IM) /3 days 8 mg (IM); Milk:36 hours 2 mg (IV/SC/IM) / 72 hours 8 mg (IM)

- **Pig (sow)**
  - Meat and offal. 4 day

- **Cattle**
  - Milk. no withdrawal period

Meat and offal: 6 days 2 mg (IV/SC/IM) /3 days 8 mg (IM); Milk:36 hours 2 mg (IV/SC/IM) / 72 hours 8 mg (IM)

Intravenous use:

- **Cattle**
  - Meat and offal. no withdrawal period

Meat and offal: 6 days 2 mg (IV/SC/IM) /3 days 8 mg (IM); Milk:36 hours 2 mg (IV/SC/IM) / 72 hours 8 mg (IM)

- **Cattle**
  - Milk. no withdrawal period

Meat and offal: 6 days 2 mg (IV/SC/IM) /3 days 8 mg (IM); Milk:36 hours 2 mg (IV/SC/IM) / 72 hours 8 mg (IM)

Subcutaneous use:

- **Cattle**

- Meat and offal. no withdrawal period

Meat and offal: 6 days 2 mg (IV/SC/IM) /3 days 8 mg (IM); Milk:36 hours 2 mg (IV/SC/IM) / 72 hours 8 mg (IM)

- **Cattle**
  - Milk. no withdrawal period

Meat and offal: 6 days 2 mg (IV/SC/IM) /3 days 8 mg (IM); Milk:36 hours 2 mg (IV/SC/IM) / 72 hours 8 mg (IM)

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA93

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

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Authorisation status:
Valid

Authorised in:
Bulgaria

Available in:
Bulgaria

Package description:
cardboard box containing 12 vials of 250 ml
cardboard box containing 12 vials of 100 ml
cardboard box containing 10 vials of 250 ml
cardboard box containing 10 vials of 100 ml
cardboard box containing 6 vials of 250 ml
cardboard box containing 6 vials of 100 ml
cardboard box containing 1 vial of 250 ml
cardboard box containing 1 vial of 100 ml

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:
Industrial Veterinaria S.A.

Marketing authorisation date:
13/02/2013

Manufacturing sites for batch release:
Industrial Veterinaria S.A.
KELA Kempisch Laboratorium Kela Laboratoria
aniMedica GmbH

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1944

Date of authorisation status change:
13/02/2013

Reference member state:
Spain

Procedure number:
ES/V/0190/001

Concerned member states:
Belgium Bulgaria Cyprus Czechia France Germany Greece Hungary Ireland Italy Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia 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: 6/04/2023
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Package Leaflet

English (PDF)
Published on: 6/04/2023
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Labelling

English (PDF)
Published on: 6/04/2023
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Combined File of all Documents