

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)

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Pig (sow)

- Meat and offal. 4 day

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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)

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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

Authorisation status:

Valid

Authorised in:

Czechia

Available in:

Czechia

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:

4/02/2014

Manufacturing sites for batch release:

Industrial Veterinaria S.A.

KELA Kempisch Laboratorium Kela Laboratoria

aniMedica GmbH

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/008/14-C

Date of authorisation status change:

4/02/2014

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)

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Combined File of all Documents