

MARBOCOLI 100 mg/ml solution for injection

Authorised

- Marbofloxacin

Product identification

Medicine name:

MARBOCOLI 100 mg/ml solution for injection

MARBOCOLI 100 mg/ml SOLUCION INYECTABLE PARA BOVINO Y PORCINO

Active substance:

Marbofloxacin

Target species:

Cattle

Pig (sow)

Pig (for fattening)

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: 3 days (8 mg (IM)). 6 days (2 mg (IV/SC/IM))

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Pig (sow)

- Meat and offal. 4 day

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Pig (for fattening)

- Meat and offal. 4 day

-

Cattle

- Milk. no withdrawal period

Milk: 72 hours (8 mg (IM)). 36 hours (2 mg (IV/SC/IM))

Intravenous use:

-

Cattle

- Meat and offal. no withdrawal period

Meat and offal: 3 days (8 mg (IM)). 6 days (2 mg (IV/SC/IM))

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Cattle

- Milk. no withdrawal period

Milk: 72 hours (8 mg (IM)). 36 hours (2 mg (IV/SC/IM))

Subcutaneous use:

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Cattle

- Meat and offal. no withdrawal period

Meat and offal: 3 days (8 mg (IM)). 6 days (2 mg (IV/SC/IM))

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Cattle

- Milk. no withdrawal period

Milk: 72 hours (8 mg (IM)). 36 hours (2 mg (IV/SC/IM))

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA93

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Available in:

Spain

Package description:

box containing 10 PP vials of 250 ml
box containing 10 PP vials of 100 ml
box containing 1 PP vial of 250 ml
box containing 1 PP vial of 100 ml
box containing 10 glass vials of 250 ml
box containing 1 glass vial of 250 ml
box containing 10 glass vials of 100 ml
box containing 1 glass vial of 100 ml
box containing 10 glass vials of 50 ml
box containing 1 glass vial of 50 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:

S P Veterinaria S.A.

Marketing authorisation date:

4/12/2014

Manufacturing sites for batch release:

S P Veterinaria S.A.

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

3151 ESP

Date of authorisation status change:

4/12/2014

Reference member state:

Spain

Procedure number:

ES/V/0217/001

Concerned member states:

Malta Poland Portugal Romania

To consult adverse reactions on veterinary medicinal products please go to

www.adrreports.eu/vet

Documents

Combined File of all Documents

Summary of Product Characteristics

English (PDF)

Published on: 11/04/2023

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Package Leaflet

English (PDF)

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Labelling

English (PDF)

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