MARBOCEN 100 mg/ml SOLUCION INYECTABLE

Authorised

- Marbofloxacin

Product identification

Medicine name:
MARBOCEN 100 mg/ml SOLUCION INYECTABLE

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

Carne:3 Días (8 mg dosis única (IM))/6 Días (2 mg entre 3 -5 días (IV/SC/IM))

  - Milk. no withdrawal period

Leche:72 Horas (8 mg dosis única (IM))/36 Horas (2 mg entre 3 -5 días (IV/SC/IM))

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

- **Pig (for fattening)**
  - Meat and offal. 4 day

Intravenous use:

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

Carne:3 Días (8 mg dosis única (IM))/6 Días (2 mg entre 3 -5 días (IV/SC/IM))

  - Milk. no withdrawal period

Leche:72 Horas (8 mg dosis única (IM))/36 Horas (2 mg entre 3 -5 días (IV/SC/IM))

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

- **Pig (for fattening)**
  - Meat and offal. 4 day

Subcutaneous use:

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

Carne:3 Días (8 mg dosis única (IM))/6 Días (2 mg entre 3 -5 días (IV/SC/IM))

  - Milk. no withdrawal period

Leche:72 Horas (8 mg dosis única (IM))/36 Horas (2 mg entre 3 -5 días (IV/SC/IM))

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

- **Pig (for fattening)**
  - Meat and offal. 4 day

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:
Spain

Package description:
Available only in Spanish
Available only in Spanish
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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:
Cenavisa S.L.

Marketing authorisation date:
20/10/2014

Manufacturing sites for batch release:
Cenavisa S.L.

Responsible authority:
Spanish Agency For Medicines And Medical Devices

Authorisation number:
3122 ESP

Date of authorisation status change:
20/10/2014

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

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

Labelling

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

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