MARBOVET 20 mg/ml SOLUCION INYECTABLE

Authorised

- Marbofloxacin

Product identification

Medicine name:
MARBOVET 20 mg/ml SOLUCION INYECTABLE

Active substance:
Marbofloxacin

Target species:
Cattle (pre-ruminant)
Cattle (calf)
Pig (for fattening)

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle (pre-ruminant)**
  - Meat and offal. 6 day

- **Cattle (calf)**
  - Meat and offal. 6 day

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

- **Cattle (pre-ruminant)**
  - Milk. no withdrawal period

leche: Su uso no está autorizado en animales cuya leche se utiliza para consumo humano

- **Cattle (calf)**
  - Milk. no withdrawal period

leche: Su uso no está autorizado en animales cuya leche se utiliza para consumo humano

Intravenous use:

- **Cattle (pre-ruminant)**
  - Meat and offal. 6 day

- **Cattle (calf)**
  - Meat and offal. 6 day

- **Cattle (pre-ruminant)**
  - Milk. no withdrawal period

leche: Su uso no está autorizado en animales cuya leche se utiliza para consumo humano

- **Cattle (calf)**
  - Milk. no withdrawal period

leche: Su uso no está autorizado en animales cuya leche se utiliza para consumo humano

Subcutaneous use:

- **Cattle (pre-ruminant)**
  - Meat and offal. 6 day

- **Cattle (calf)**
  - Meat and offal. 6 day

- **Cattle (pre-ruminant)**
  - Milk. no withdrawal period

leche: Su uso no está autorizado en animales cuya leche se utiliza para consumo humano

- **Cattle (calf)**
  - Milk. no withdrawal period

leche: Su uso no está autorizado en animales cuya leche se utiliza para consumo humano

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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:
Available only in Spanish
Available only in Spanish
Available only in Spanish

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:
Fatro S.p.A.

Marketing authorisation date:
2/02/2012

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Spanish Agency For Medicines And Medical Devices

Authorisation number:
2450 ESP

Date of authorisation status change:
2/02/2012

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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