MARBIFLOX 100 MG/ML SOLUTION FOR INJECTION FOR BOVINES AND PIGS (SOWS)

Authorised

- Marbofloxacin

Product identification

Medicine name:
MARBIFLOX 100 MG/ML SOLUTION FOR INJECTION FOR BOVINES AND PIGS (SOWS)

Active substance:
Marbofloxacin

Target species:
Cattle
Pig (sow)

Route of administration:
Intramuscular 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. 3 day — 8 mg/kg single dose
  - Milk. 36 hour — 2 mg/kg single daily injection for 3 days
  - Meat and offal. 6 day — 2 mg/kg single daily injection for 3 days
  - Milk. 72 hour — 8 mg/kg single dose
- **Pig (sow)**
  - Meat and offal. 4 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 6 day — 2 mg/kg single daily injection for 3 days
  - Milk. 36 hour — 2 mg/kg single daily injection for 3 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA93

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

Authorisation status:
Valid

Authorised in:
France

Package description:
Box of one 50 ml solution bottle for injection
Box of one 250 ml solution bottle for injection
Box of one 100 ml solution bottle for injection

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:
KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:
4/02/2011

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto
Virbac

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/9951358 0/2011

Date of authorisation status change:
5/07/2016

Reference member state:
France

Procedure number:
FR/V/0222/002

Concerned member states:

Germany Portugal United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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

Combined File of all Documents

Package Leaflet and Labelling

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