File downloaded on 2026-11-08
Source URL: https://medicines.health.europa.eu/veterinary/en/600000046422

Marfloxin 20 mg/ml solution for injection for calves, pigs, dogs and cats

Not authorised

- Marbofloxacin

Product identification

Medicine name:
Marfloxin 20 mg/ml solution for injection for calves, pigs, dogs and cats

Active substance:
Marbofloxacin

Target species:
Cattle
Pig
Dog
Cat

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**
  - Meat and offal. 4 day

- **Pig**
  - Meat and offal. 2 day

Intravenous use:

- **Cattle**
  - Meat and offal. 4 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA93

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

---

Authorisation status:

Surrendered

Authorised in:
Slovenia

Package description:
20 ml amber glass vial (Ph. Eur. type II) sealed with a bromobutyl rubber stopper and aluminium closure packaged in an outer carton.
50 ml amber glass vial (Ph. Eur. type II) sealed with a bromobutyl rubber stopper and aluminium closure packaged in an outer carton.
100 ml amber glass vial (Ph. Eur. type II) sealed with a bromobutyl rubber stopper and aluminium closure packaged in an outer carton.

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:
17/05/2011

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

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0426/001

Date of authorisation status change:
24/01/2023

Reference member state:

Ireland

Procedure number:
IE/V/0262/001

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 3/05/2024
Updated on: 14/03/2026
Download

Package Leaflet and Labelling

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