ENROXIL 50 mg/ml solution for injection for calves, pigs and dogs

Authorised

- Enrofloxacin

Product identification

Medicine name:
ENROXIL 50 mg/ml solution for injection for calves, pigs and dogs

Active substance:
Enrofloxacin

Target species:
Pig
Cattle
Dog

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Enrofloxacin
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - Meat and offal. 13 day

Intravenous use:

- **Cattle**
  - Meat and offal. 5 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 12 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

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

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Cardboard box with 1 amber Type II glass bottle of 100 ml with a grey bromobutyl rubber stopper and aluminiumcap.
Cardboard box with 1 amber Type II glass bottle of 50 ml with a grey bromobutyl rubber stopper and aluminium cap.
Cardboard box with 1 amber Type I glass bottle of 100 ml with a grey bromobutyl rubber stopper and aluminiumcap.

Cardboard box with 1 amber Type I glass bottle of 50 ml with a grey bromobutyl rubber stopper and aluminiumcap.

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:
15/03/2011

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

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 10487

Date of authorisation status change:
30/06/2022

Reference member state:
Ireland

Procedure number:
IE/V/0422/001

Concerned member states:
Austria Germany Netherlands United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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

Summary of Product Characteristics