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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:
United Kingdom (Northern Ireland)

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:
15/03/2006

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 01656/3068

Date of authorisation status change:
11/05/2024

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

Summary of Product Characteristics