

Enroxil 100 mg/ml Solution for Injection for Cattle and Pigs

Authorised

- Enrofloxacin

Product identification

Medicine name:

Enroxil 100 mg/ml Solution for Injection for Cattle and Pigs

Active substance:

Enrofloxacin

Target species:

Pig

Cattle

Route of administration:

Intramuscular use

Intravenous use

Subcutaneous use

Product details

Active substance and strength:

Enrofloxacin

100.00 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

- Milk. 3 day

Subcutaneous use:

-

Cattle

- Meat and offal. 12 day

- Milk. 4 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 I glass bottle of 100 ml with a grey bromobutyl rubber stopper and aluminium cap.

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

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 II glass bottle of 100 ml with a grey bromobutyl rubber stopper and aluminium cap.

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:

KRKA tovarna zdravil d.d. Novo mesto

TAD Pharma GmbH

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 01656/3067

Date of authorisation status change:

18/05/2024

Reference member state:

Ireland

Procedure number:

IE/V/0422/002

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