ENRODEXIL 100 mg/ml solution for injection for cattle and pigs

Authorised

- Enrofloxacin

Product identification

Medicine name:
ENRODEXIL 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. 72 hour

Subcutaneous use:

- **Cattle**
  - Meat and offal. 12 day
  - Milk. 96 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

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

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:

Type II Amber glass vial of 100 ml capacity closed with grey bromobutyl rubber stopper and aluminium flip-offseal. One vial of 100 ml is available in a cardboard box. Type II Amber glass vial of 250 ml capacity closed with pink bromobutyl rubber stopper and aluminium flip-offseal. One vial of 250 ml is available in a cardboard box.

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:
Industrial Veterinaria S.A.

Marketing authorisation date:
23/05/2011

Manufacturing sites for batch release:
Industrial Veterinaria S.A.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V392962

Date of authorisation status change:
23/05/2011

Reference member state:
Ireland

Procedure number:
IE/V/0264/001

Concerned member states:
Austria Belgium Estonia Germany Greece Hungary Italy Netherlands Poland

Portugal Romania Spain 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.

Package Leaflet

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

Labelling

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