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

Authorised

- Enrofloxacin

Product identification

Medicine name:
Enrocare 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:
Ireland

Package description:
100 ml Amber Type I multidose glass vial with a grey bromobutyl rubber stopper and aluminium overseal.100 ml vial sold in packs containing 1 x 100 ml cartons.

100 ml Amber Type I multidose glass vial with a grey bromobutyl rubber stopper and aluminium overseal.100 ml vial sold in packs containing 6 x 100 ml cartons.
100 ml Amber Type I multidose glass vial with a grey bromobutyl rubber stopper and aluminium overseal.100 ml vial sold in packs containing 12 x 100 ml cartons.
250 ml Amber Type I multidose glass vial with a grey bromobutyl rubber stopper and aluminium overseal.250 ml vial sold in packs containing 1 x 250 ml cartons.
250 ml Amber Type I multidose glass vial with a grey bromobutyl rubber stopper and aluminium overseal.250 ml vial sold in packs containing 6 x 250 ml cartons.
250 ml Amber Type I multidose glass vial with a grey bromobutyl rubber stopper and aluminium overseal.250 ml vial sold in packs containing 12 x 250 ml cartons.

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:
Emdoka

Marketing authorisation date:
27/09/2013

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10534/007/003

Date of authorisation status change:
27/09/2013

Reference member state:
Ireland

Procedure number:
IE/V/0455/003

Concerned member states:
Luxembourg 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