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

Authorised

- Enrofloxacin

Product identification

Medicine name:
Unisol 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

Available in:
Ireland

Package description:

Type II Amber glass vial of 100 ml capacity closed with grey bromobutyl rubber stopper and aluminium flip-off seal.Cardboard box containing 1 vial of 100 ml
Type II Amber glass vial of 100 ml capacity closed with grey bromobutyl rubber stopper and aluminium flip-off seal.Cardboard box containing 12 vials of 100 ml
Type II Amber glass vial of 250 ml capacity closed with pink bromobutyl rubber stopper and aluminium flip-off seal.Cardboard box containing 1 vial of 250 ml
Type II Amber glass vial of 250 ml capacity closed with pink bromobutyl rubber stopper and aluminium flip-off seal.Cardboard box containing 12 vials of 250 ml.

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:
Vetpharma Animal Health S.L.

Marketing authorisation date:
6/05/2011

Manufacturing sites for batch release:
Industrial Veterinaria S.A.
Chemical Iberica Productos Veterinarios S.L.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10516/001/001

Date of authorisation status change:
6/05/2011

Reference member state:
Ireland

Procedure number:
IE/V/0255/001

Concerned member states:
Austria Cyprus Germany Hungary Italy Poland Portugal 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