

Enrotron 100 mg/ml Solution for injection for cattle, sheep, goats and pigs

Authorised

- Enrofloxacin

Product identification

Medicine name:

Enrotron 100 mg/ml Solution for injection for cattle, sheep, goats and pigs

Active substance:

Enrofloxacin

Target species:

Pig
Cattle
Goat
Sheep

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

-

Goat

- Meat and offal. 6 day

- Milk. 4 day

-

Sheep

- Meat and offal. 4 day

- Milk. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Available in:

Italy

Package description:

Pack Size: 100 ml clear glass vial type I with siliconised, Teflon coated rubber stopper sealed with an aluminium cap. Carton of 1 x 100 ml is available.

Pack Size: 100 ml clear glass vial type I with siliconised, Teflon coated rubber stopper sealed with an aluminium cap. Cartons of 12 x 100 ml are available.

Pack Size: 100 ml clear glass vial type I with siliconised, uncoated bromobutyl rubber stopper, sealed with an aluminium cap. Carton of 1 x 100 ml is available.

Pack Size: 100 ml clear glass vial type I with siliconised, uncoated bromobutyl rubber stopper, sealed with an aluminium cap. Cartons of 12 x 100 ml are available.

Pack Size: 250 ml clear glass vial type I with siliconised, uncoated bromobutyl rubber stopper, sealed with an aluminium cap. Carton of 1 x 250 ml is available.

Pack Size: 250 ml clear glass vial type I with siliconised, uncoated bromobutyl rubber stopper, sealed with an aluminium cap. Cartons of 12 x 250 ml are available.

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:

aniMedica GmbH

Marketing authorisation date:

1/06/2012

Manufacturing sites for batch release:

Industrial Veterinaria S.A.

aniMedica GmbH

Responsible authority:

Ministry Of Health

Authorisation number:

104370

Date of authorisation status change:

1/04/2018

Reference member state:

Ireland

Procedure number:

IE/V/0270/003

Concerned member states:

Estonia Greece Hungary Italy Latvia Lithuania Poland Portugal Romania
Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to

www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

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Summary of Product Characteristics