

Valemas 50 mg/ml Solution for Injection for Cattle, Sheep, Goats, Pigs, Dogs and Cats

Authorised

- Enrofloxacin

Product identification

Medicine name:

Valemas 50 mg/ml Solution for Injection for Cattle, Sheep, Goats, Pigs, Dogs and Cats

Active substance:

Enrofloxacin

Target species:

Pig
Cattle
Dog
Goat
Sheep
Cat

Route of administration:

Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

Enrofloxacin

50.00 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

Subcutaneous use:

-

Cattle

- Meat and offal. 12 day

-

Goat

- Meat and offal. 6 day

- Milk. 96 hour

-

Sheep

- Meat and offal. 4 day

- Milk. 72 hour

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)

Available in:

United Kingdom (Northern Ireland)

Package description:

The product is presented in amber Type II glass bottle of 50 ml with chlorobutyl rubber stopper sealed with a flip-off aluminium cap with a tamper-evident polypropylene seal.

The product is presented in amber Type II glass bottle of 100 ml with chlorobutyl rubber stoppers sealed with a flip-off aluminium cap with a tamper-evident polypropylene seal.

The product is presented in amber Type II glass bottle of 250 ml with chlorobutyl rubber stoppers sealed with a flip-off aluminium cap with a tamper-evident polypropylene seal.

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:

Fatro S.p.A.

Marketing authorisation date:

21/05/2019

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 11557/4003

Date of authorisation status change:

19/11/2019

Reference member state:

Ireland

Procedure number:

IE/V/0445/001

Concerned member states:

United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics