

# 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](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics