

# HEMOSILATE 125 mg/ml solution for injection

Authorised

- Etamsylate

## Product identification

**Medicine name:**

HEMOSILATE 125 mg/ml solution for injection

**Active substance:**

Etamsylate

**Target species:**

Cattle

Sheep

Goat

Pig

Horse

Dog

Cat

**Route of administration:**

Intravenous use

Intramuscular use

## Product details

### **Active substance and strength:**

Etamsylate

125.00 milligram(s) / 1.00 millilitre(s)

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### **Pharmaceutical form:**

Solution for injection

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### **Withdrawal period by route of administration:**

#### **Intravenous use:**

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#### **Cattle**

- Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- Milk. 0 day

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#### **Sheep**

- Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- Milk. 0 day

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#### **Goat**

- Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- Milk. 0 day

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#### **Pig**

- Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

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### **Horse**

- Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- Milk. 0 day

### **Intramuscular use:**

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### **Cattle**

- Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- Milk. 0 day

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### **Sheep**

- Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- Milk. 0 day

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### **Goat**

- Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- Milk. 0 day

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### **Pig**

- Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

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### **Horse**

- Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- Milk. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QB02BX01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Malta

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**Package description:**

cardboard box containing 10 vials of 20 ml

cardboard box containing 5 vials of 20 ml

cardboard box containing 1 vial of 20 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ecuphar Veterinaria S.L.U.

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**Marketing authorisation date:**

6/03/2018

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**Manufacturing sites for batch release:**

Zoetis Manufacturing & Research Spain S.L.

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**Responsible authority:**

Ministry For Agriculture Fisheries And Animal Rights

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**Authorisation number:**

VMA68

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**Date of authorisation status change:**

6/03/2018

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0281/001

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**Concerned member states:**

Austria Belgium Cyprus Czechia Denmark Estonia Finland France Germany  
Greece Hungary Ireland Italy Malta Netherlands Norway Poland Portugal  
Romania Slovakia Slovenia Sweden United Kingdom (Northern Ireland)

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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

Package Leaflet

eu-PUAR-esv0281001-dcp-hemosilate-125-mg-ml-solution-for-injection-en.pdf