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# HEMOSILATE 125 mg/ml solution for injection

Authorised

- Etamsilate

## Product identification

**Medicine name:**
HEMOSILATE 125 mg/ml solution for injection

**Active substance:**
Etamsilate

**Target species:**
Cattle
Sheep
Goat
Pig
Horse
Dog
Cat

**Route of administration:**
Intravenous use
Intramuscular use

## Product details

**Active substance and strength:**
Etamsilate
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:**

- **Cattle**
  - Meat and offal. no withdrawal period

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

  - Milk. 0 day

- **Sheep**
  - Meat and offal. no withdrawal period

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

  - Milk. 0 day

- **Goat**
  - Meat and offal. no withdrawal period

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

  - Milk. 0 day

- **Pig**
  - Meat and offal. no withdrawal period

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

- **Horse**
  - Meat and offal. no withdrawal period

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

  - Milk. 0 day

**Intramuscular use:**

- **Cattle**
  - Meat and offal. no withdrawal period

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

  - Milk. 0 day

- **Sheep**
  - Meat and offal. no withdrawal period

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

  - Milk. 0 day

- **Goat**
  - Meat and offal. no withdrawal period

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

  - Milk. 0 day

- **Pig**
  - Meat and offal. no withdrawal period

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

- **Horse**

- Meat and offal. no withdrawal period

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

- Milk. 0 day

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QB02BX01

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Sweden

**Package description:**
Available only in Swedish
Available only in Swedish
Available only in Swedish

## 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:**
Ecuphar Veterinaria S.L.U.

**Marketing authorisation date:**
26/05/2020

**Manufacturing sites for batch release:**
Zoetis Manufacturing & Research Spain S.L.

**Responsible authority:**
Swedish Medical Products Agency

**Authorisation number:**
58777

**Date of authorisation status change:**
26/05/2020

**Reference member state:**
Spain

**Procedure number:**
ES/V/0281/001

**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)

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

## Documents

Summary of Product Characteristics

English (PDF)
Published on: 14/03/2026
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Package Leaflet

English (PDF)
Published on: 14/03/2026
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Labelling

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