# E-SELENSOL 70/1 mg/ml emulsion for injection for cattle, sheep and pigs

Authorised

- ALPHA-TOCOPHEROL
- Sodium selenite

## Product identification

**Medicine name:**
E-SELENSOL 70/1 mg/ml emulsion for injection for cattle, sheep and pigs
E-Selensol, 70/1 mg/ml süsteemulsioon veistele, lammastele ja sigadele

**Active substance:**
ALPHA-TOCOPHEROL
Sodium selenite

**Target species:**
Cattle
Pig
Sheep

**Route of administration:**
Intramuscular use
Subcutaneous use

## Product details

**Active substance and strength:**
ALPHA-TOCOPHEROL
70.00 milligram(s) / 1.00 millilitre(s)

Sodium selenite
1.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**
Emulsion for injection

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**Withdrawal period by route of administration:**
**Intramuscular use:**

- **Cattle**
  - Meat and offal. 14 day

Not authorised for use in animals producing milk for human consumption.

- **Pig**
  - Meat and offal. 14 day

- **Sheep**
  - Meat and offal. 30 day

Not authorised for use in animals producing milk for human consumption.

**Subcutaneous use:**

- **Cattle**
  - Meat and offal. 14 day

Not authorised for use in animals producing milk for human consumption.

- **Pig**

- Meat and offal. 14 day

- **Sheep**
  - Meat and offal. 30 day

Not authorised for use in animals producing milk for human consumption.

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QA12CE99

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

**Authorisation status:**
Valid

**Authorised in:**
Estonia

**Available in:**
Estonia

**Package description:**
Type II amber coloured glass vials, with bromobutyl rubber stopper and lacquered aluminium cap. Carton box with 1 vial of 50 ml.
Type II amber coloured glass vials, with bromobutyl rubber stopper and lacquered aluminium cap. Carton box with 1 vial of 100 ml

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Generic (abridged application) - art 13(1)

**Marketing authorisation holder:**
Labiana Life Sciences S.A.

**Marketing authorisation date:**
27/04/2021

**Manufacturing sites for batch release:**
Labiana Life Sciences S.A.

**Responsible authority:**
State Agency Of Medicines

**Authorisation number:**
2285

**Date of authorisation status change:**
27/04/2021

**Reference member state:**
Hungary

**Procedure number:**
HU/V/0143/001

**Concerned member states:**
Croatia Cyprus Estonia Greece Latvia Lithuania

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

## Documents

Summary of Product Characteristics

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