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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

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)

Pharmaceutical form:
Emulsion for injection

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

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.