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)

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

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Authorisation status:
Valid

Authorised 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:
28/04/2021

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

Responsible authority:
State Agency Of Medicines

Authorisation number:
2285

Date of authorisation status change:
28/04/2021

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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