

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.

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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

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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:

Croatia

Available in:

Croatia

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:

5/05/2021

Manufacturing sites for batch release:

Labiana Life Sciences S.A.

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/21-01/283

Date of authorisation status change:

17/08/2021

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