

LABIASOL E SELEN, (70+2,3)MG/1ML ενέσιμο διαλύμα για βοοειδή, πρόβατα και χοίρους

Authorised

- Sodium selenite
- TOCOPHERYL ACETATE

Product identification

Medicine name:

LABIASOL E SELEN, (70+2,3)MG/1ML ενέσιμο διαλύμα για βοοειδή, πρόβατα και χοίρους

Active substance:

Sodium selenite

TOCOPHERYL ACETATE

Target species:

Cattle

Sheep

Pig

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Sodium selenite

2.30 milligram(s) / 1.00 millilitre(s)

TOCOPHERYL ACETATE

70.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Meat and offal. 30 day

-

Sheep

- Meat and offal. 8 day

-

Pig

- Meat and offal. 8 day

Subcutaneous use:

-

Cattle

- Meat and offal. 30 day

-

Sheep

- Meat and offal. 8 day

-

Pig

- Meat and offal. 8 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12CE99

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Package description:

Available only in [Greek](#)

Available only in [Greek](#)

Available only in [Greek](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone)

Marketing authorisation holder:

Chellafarm Vet A.E.

Marketing authorisation date:

7/11/1994

Manufacturing sites for batch release:

Labiana Life Sciences S.A.

Responsible authority:

National Organization For Medicines

Authorisation number:

34728/08-11-1994/K-0088201

Date of authorisation status change:

25/08/2020

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