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

Ima dovoljenje za promet

- ALPHA-TOCOPHEROL
- Sodium selenite

Informacije o zdravilu

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

Učinkovina:
Na voljo samo v English
Na voljo samo v English

Ciljne živalske vrste:
govedo
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Pot uporabe:
intramuskularna uporaba
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
70.00 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
1.00 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:

Emulzija za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

- **govedo**
  - 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.

Subkutana uporaba:

- **govedo**
  - 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.

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QA12CE99

Pravni status za dobavo/izdajo zdravila:
Zdravilo, ki se izdaja na veterinarski recept

Status dovoljenja za promet z zdravilom:
Valid

Ima dovoljenje za promet v:
Na voljo samo v Spanish German Estonian English French Italian Dutch Portuguese Slovak Finnish Swedish Icelandic Norwegian

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Na voljo v:
Estonia

Opis ovojnine zdravila:
Na voljo samo v English
Na voljo samo v English

Dodatne informacije

Vrsta dovoljenja:
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

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Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English

Imetnik dovoljenja za promet z zdravilom:

Labiana Life Sciences S.A.

Datum dovoljenja za promet z zdravilom:
27/04/2021

Proizvodna mesta, odgovorna za sproščanje serij:
Labiana Life Sciences S.A.

Pristojni organ:
State Agency Of Medicines

Številka dovoljenja za promet z zdravilom:
2285

Datum spremembe statusa dovoljenja za promet:
27/04/2021

Referenčna država članica:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Številka postopka:
HU/V/0143/001

Zadevne države članice:
Na voljo samo v Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian Greek English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish German Estonian English French Italian Dutch Portuguese Slovak Finnish Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Povzetek glavnih značilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Navodilo za uporabo

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Označevanje

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