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NEOCAL 300

Ima dovoljenje za promet

- Caffeine
- Magnesium chloride hexahydrate
- CALCIUM GLYCEROPHOSPHATE
- Calcium gluconate

Informacije o zdravilu

Ime zdravila:
NEOCAL 300

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

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

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
20.00 milligram(s) / 1.00 millilitre(s)

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

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

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

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:
Subkutana uporaba:

- **Horse**
  - Meat and offal. 0 day
  - Мляко: нула дни.
-

govedo

- Meat and offal. 0 day

Мляко: нула дни.

- **Sheep**
  - Meat and offal. 0 day

Мляко: нула дни.

- **Goat**
  - Meat and offal. 0 day

Мляко: нула дни.

- **Pig**
  - Meat and offal. 0 day

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

QA12AX

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 Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Opis ovojnine zdravila:

Na voljo samo v Bulgarian

Na voljo samo v Bulgarian

Dodatne informacije

Vrsta dovoljenja:

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

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

Imetnik dovoljenja za promet z zdravilom:
Dsm Denitrans OOD

Datum dovoljenja za promet z zdravilom:
20/03/2019

Proizvodna mesta, odgovorna za sproščanje serij:
Dsm Denitrans OOD

Pristojni organ:
Bulgarian Food Safety Authority

Številka dovoljenja za promet z zdravilom:
0022-2872

Datum spremembe statusa dovoljenja za promet:
25/10/2022

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.