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DEVOMYCIN, 200MG/ML ενέσιμο διάλυμα για βοοειδή, πρόβατα, χοίρους, σκύλους και γάτες

Ima dovoljenje za promet

- Streptomycin sulfate

Informacije o zdravilu

Ime zdravila:
DEVOMYCIN, 200MG/ML ενέσιμο διάλυμα για βοοειδή, πρόβατα, χοίρους, σκύλους και γάτες

Učinkovina:
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
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

Podatki o zdravilu

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

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **govedo**
  - Meat and offal. 16 day
  - Milk. 48 hour
- **Sheep**
  - Meat and offal. 19 day
  - Milk. 48 hour
- **Pig**
  - Meat and offal. 19 day
- **Dog**
  - Meat and offal. no withdrawal period Δεν εφαρμόζεται
- **Cat**

- Meat and offal. no withdrawal period Δεν εφαρμόζεται

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QJ01GA01

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

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

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 Portuguese

Imetnik dovoljenja za promet z zdravilom:
Hellafarm S.A.

Datum dovoljenja za promet z zdravilom:
21/07/1988

Proizvodna mesta, odgovorna za sproščanje serij:
Norbrook Laboratories (Ireland) Limited

Pristojni organ:
National Organization For Medicines

Številka dovoljenja za promet z zdravilom:
51833/13-07-2012/K-0029001

Datum spremembe statusa dovoljenja za promet:
25/08/2020

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