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TYLUCYL 200 MG/ML SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Ima dovoljenje za promet

- Tylosin

Informacije o zdravilu

Ime zdravila:
TYLUCYL 200 MG/ML SOLUTION FOR INJECTION FOR CATTLE AND PIGS

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

Pot uporabe:
intramuskularna uporaba
Intravenska uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
200000.00 international unit(s) / 1.00 millilitre(s)

Farmacevtska oblika:

Raztopina za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

- **govedo**
  - Meat and offal. 28 day
  - Milk. 108 hour
- **Pig**
  - Meat and offal. 16 day

Intravenska uporaba:

- **govedo**
  - Meat and offal. 28 day
  - Milk. 108 hour

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

QJ01FA90

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 French
Na voljo samo v French
Na voljo samo v French

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:
Vetoquinol Biowet Sp. z o.o.

Datum dovoljenja za promet z zdravilom:
17/11/2016

Proizvodna mesta, odgovorna za sproščanje serij:
Vetoquinol S.A.

Pristojni organ:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Številka dovoljenja za promet z zdravilom:
2579

Datum spremembe statusa dovoljenja za promet:
17/11/2016

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

Številka postopka:
FR/V/0289/001

Zadevne države članice:

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

Dokumenti

Navodilo za uporabo

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

Povzetek glavnih značilnosti zdravila

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

Označevanje

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Combined File of all Documents

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