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Tardocyclin Spray, 23,34 mg/g, sprej za kožu, suspenzija, za konja, govedo, ovcu, kozu, svinju, psa i mačku

Ima dovoljenje za promet

- Oxytetracycline hydrochloride

Informacije o zdravilu

Ime zdravila:
Tardocyclin Spray, 23,34 mg/g, sprej za kožu, suspenzija, za konja, govedo, ovcu, kozu, svinju, psa i mačku

Učinkovina:
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
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:
Dermalna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
23.34 milligram(s) / 1.00 gram(s)

Farmacevtska oblika:
Dermalno pršilo, suspenzija

Karenca glede na pot uporabe:
Dermalna uporaba:

- **Horse**
  - Meat and offal. 0 day
- **govedo**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 0 day

- Milk. 0 day

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

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

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QD06AA03

Pravni status za dobavo/izdajo zdravila:
Zdravilo, ki se izdaja brez veterinarskega recepta

Status dovoljenja za promet z zdravilom:
Valid

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

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Opis ovojnine zdravila:
Na voljo samo v Croatian

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 Italian

Imetnik dovoljenja za promet z zdravilom:

Bayer d.o.o.

Datum dovoljenja za promet z zdravilom:
7/07/2015

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

Pristojni organ:
Croatian Veterinary Institute

Številka dovoljenja za promet z zdravilom:
UP/I-322-05/13-01/489

Datum spremembe statusa dovoljenja za promet:
7/07/2015

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.