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Cyclospray 78.6 mg/g, cutaneous spray, suspension for pigs, sheep and cattle

Ima dovoljenje za promet

- Chlortetracycline hydrochloride

Informacije o zdravilu

Ime zdravila:
Cyclospray 78.6 mg/g, cutaneous spray, suspension for pigs, sheep and cattle

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

Pot uporabe:
Dermalna uporaba

Podatki o zdravilu

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

Farmacevtska oblika:
Dermalno pršilo, raztopina

Karenca glede na pot uporabe:
Dermalna uporaba:

- **govedo**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Pig**
  - Meat and offal. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QD06AA02

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 Lithuanian Dutch Portuguese Romanian Slovak Swedish Icelandic Norwegian

Na voljo v:
Romania

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

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

Imetnik dovoljenja za promet z zdravilom:
Eurovet Animal Health B.V.

Datum dovoljenja za promet z zdravilom:
17/01/2016

Proizvodna mesta, odgovorna za sproščanje serij:
IGS Aerosols GmbH
Eurovet Animal Health B.V.

Pristojni organ:
Institute For Control Of Biological Products And Veterinary Medicines

Številka dovoljenja za promet z zdravilom:
200153

Datum spremembe statusa dovoljenja za promet:
19/02/2024

Referenčna država članica:

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

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Številka postopka:
PT/V/0132/001

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 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
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Na voljo samo v Estonian English French Lithuanian Portuguese 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

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.