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Ophtocycline 10 mg/g mazilo za oko za pse, mačke in konje

Ima dovoljenje za promet

- Chlortetracycline hydrochloride

Informacije o zdravilu

Ime zdravila:
Ophtocycline 10 mg/g mazilo za oko za pse, mačke in konje

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
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:
Okularna uporaba

Podatki o zdravilu

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

Farmacevtska oblika:
Mazilo za oko

Karenca glede na pot uporabe:
Okularna uporaba:

- **Horse**
  - Meat and offal. 1 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QS01AA02

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

Na voljo v:
Slovenia

Opis ovojnine zdravila:
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:
Le Vet. Beheer B.V.

Datum dovoljenja za promet z zdravilom:
2/10/2017

Proizvodna mesta, odgovorna za sproščanje serij:
Produlab Pharma B.V.

Pristojni organ:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Številka dovoljenja za promet z zdravilom:
DC/V/0586/001

Datum spremembe statusa dovoljenja za promet:
2/10/2017

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

Številka postopka:
NL/V/0209/001

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
Na voljo samo v Spanish Czech German Estonian 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

Navodilo za uporabo

Označevanje

Combined File of all Documents

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