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Cefenicol CA 5 mg/ml eye drops, solution for dogs and cats

Ima dovoljenje za promet

- Chloramphenicol

Informacije o zdravilu

Ime zdravila:
Cefenicol CA 5 mg/ml eye drops, solution for dogs and cats

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

Pot uporabe:
Okularna uporaba

Podatki o zdravilu

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

Farmacevtska oblika:
Kapljice za oko, raztopina

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QS01AA01

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:
Germany

Opis ovojnine zdravila:
Na voljo samo v German

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 French Italian Latvian Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
CP-Pharma Handelsgesellschaft mbH

Datum dovoljenja za promet z zdravilom:
27/12/2018

Proizvodna mesta, odgovorna za sproščanje serij:
CP-Pharma Handelsgesellschaft mbH

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
402485.00.00

Datum spremembe statusa dovoljenja za promet:
27/12/2018

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

Številka postopka:
DE/V/0344/001

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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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

Combined File of all Documents

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

2402485-paren-20240710.pdf

2402485-parde-20181214.rtf