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Neptra 16.7 mg/ml florfenicol + 16.7 mg/ml terbinafine hydrochloride + 2.2 mg/ml mometasone furoate - Ear drops, solution in single-dose container

Ima dovoljenje za promet

- Florfenicol
- Terbinafine hydrochloride
- Mometasone furoate

Informacije o zdravilu

Ime zdravila:
Neptra 16.7 mg/ml florfenicol + 16.7 mg/ml terbinafine hydrochloride + 2.2 mg/ml mometasone furoate - Ear drops, solution in single-dose container

Učinkovina:
Na voljo samo v English
Na voljo samo v English
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

Pot uporabe:
Avrikularna uporaba

Podatki o zdravilu

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

Na voljo samo v English
16.70 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
2.20 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Kapljice za uho, raztopina v enoodmernem vsebniku

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QS02CA91

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
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Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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Na voljo v:
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Opis ovojnine zdravila:

Na voljo samo v English

Na voljo samo v English
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:
Elanco Animal Health GmbH

Datum dovoljenja za promet z zdravilom:
10/12/2019

Proizvodna mesta, odgovorna za sproščanje serij:
KVP Pharma+Veterinaer Produkte GmbH

Pristojni organ:
European Commission

Številka dovoljenja za promet z zdravilom:
Ta podatek za to zdravilo ni na voljo.

Datum spremembe statusa dovoljenja za promet:
10/12/2019

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

Dokumenti

Combined File of all Documents

slovenščina (PDF)
Objavljeno na dan: 13/07/2026
Prenesi

ema-puar-neptra-v-4735-par-en.pdf