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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

Atļautas

- Florfenicol
- Terbinafine hydrochloride
- Mometasone furoate

Zāļu identifikācija

Zāļu nosaukums:
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

Aktīvā viela:
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English

Mērķsugas:
Suns

Lietošanas veids:
Lietošanai ausī

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:

Pieejamas tikai English
16.70 miligrams(i) / 1.00 mililitrs(i)

Pieejamas tikai English
16.70 miligrams(i) / 1.00 mililitrs(i)

Pieejamas tikai English
2.20 miligrams(i) / 1.00 mililitrs(i)

Farmaceitiskā forma:

Ausu pilieni, šķīdums vienas devas iepakojumā

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:

QS02CA91

Izplatīšanas juridiskie nosacījumi:

Recepšu veterinārās zāles

Atļaujas statuss:

Derīga

Atļautas:

Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Pieejamas tikai Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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Pieejams:

Austria , Belgium , Bulgaria , Croatia , Cyprus , 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) , Čehija

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Iepakojuma apraksts:

Pieejamas tikai English

Pieejamas tikai English

Pieejamas tikai English

Pieejamas tikai English

Papildu informācija

Tiesību tips:

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:
Pieteikums fiksētas kombinācijas veterinārajām zālēm (Direktīvas 2001/82/EC 13.(b) pants)

Tirdzniecības atļaujas turētājs:
Elanco Animal Health GmbH

Tirdzniecības atļaujas datums:
10/12/2019

Ražošanas vietas sēriju izlaidei:
KVP Pharma+Veterinaer Produkte GmbH

Atbildīgā iestāde:
European Commission

Atļaujas numurs:
Šī informācija par šīm zālēm nav pieejama.

Atļaujas statusa maiņas datums:
10/12/2019

Lai uzzinātu par veterināro zāļu blakusparādībām, lūdzu, dodieties uz: www.adrreports.eu/vet

Dokumenti

Visu dokumentu apvienotais fails

latviešu (PDF)

Publicēts: 13/07/2026

Lejupielādēt

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