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Otoxolan ear drops, suspension for dogs

Ima dovoljenje za promet

- Marbofloxacin
- Clotrimazole
- Dexamethasone acetate

Informacije o zdravilu

Ime zdravila:
Otoxolan ear drops, suspension for dogs

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
3.00 milligram(s) / 1.00 millilitre(s)

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

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

Farmacevtska oblika:

Kapljice za uho, suspenzija

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

QS02CA06

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

Opis ovojnine zdravila:

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:

KRKA tovarna zdravil d.d. Novo mesto

Datum dovoljenja za promet z zdravilom:
18/11/2016

Proizvodna mesta, odgovorna za sproščanje serij:
TAD Pharma GmbH
KRKA tovarna zdravil d.d. Novo mesto

Pristojni organ:
French Agency For Food, Environmental And Occupational Health & Safety

Številka dovoljenja za promet z zdravilom:
FR/V/8243893 5/2016

Datum spremembe statusa dovoljenja za promet:
22/12/2021

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

Številka postopka:
IE/V/0438/001

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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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.

Package Leaflet and Labelling

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