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Ototop Ear Drops and Cutaneous Suspension for Dogs, Cats and Guinea Pigs

Ima dovoljenje za promet

- POLYMYXIN B SULFATE
- Prednisolone acetate
- Miconazole nitrate

Informacije o zdravilu

Ime zdravila:
Ototop Ear Drops and Cutaneous Suspension for Dogs, Cats and Guinea Pigs

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
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:

Dermalna uporaba

Avrikularna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
5500.00 international unit(s) / 1.00 millilitre(s)

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

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

Farmacevtska oblika:

Na voljo samo v Bulgarian Spanish Czech Danish Greek English Irish Latvian Lithuanian Romanian Slovak Finnish Swedish

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

QS02CA01

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 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:
Industrial Veterinaria S.A.

Datum dovoljenja za promet z zdravilom:
3/04/2020

Proizvodna mesta, odgovorna za sproščanje serij:
Industrial Veterinaria S.A.
aniMedica Herstellungs GmbH
aniMedica GmbH

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
402648.00.00

Datum spremembe statusa dovoljenja za promet:
3/04/2020

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

Številka postopka:
DE/V/0321/001

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 Greek English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Na voljo samo v Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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.

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