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Mitex Ohrentropfen und Suspension zur Anwendung auf der Haut für Hunde und Katzen

Ima dovoljenje za promet

- Miconazole nitrate
- Prednisolone acetate
- POLYMYXIN B SULFATE

Informacije o zdravilu

Ime zdravila:
Mitex Ohrentropfen und Suspension zur Anwendung auf der Haut für Hunde und Katzen

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

Pot uporabe:

Avrikularna uporaba
Dermalna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

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

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

Na voljo samo v English
5500.00 international unit(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:

Denmark

Opis ovojnine zdravila:

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:
Vetviva Richter GmbH

Datum dovoljenja za promet z zdravilom:
12/03/2015

Proizvodna mesta, odgovorna za sproščanje serij:
Vetviva Richter GmbH

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
53793

Datum spremembe statusa dovoljenja za promet:
12/03/2015

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

Številka postopka:
AT/V/0014/001

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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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.

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