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OTOMAX EAR DROPS SUSPENSION

Ima dovoljenje za promet

- Gentamicin sulfate
- Clotrimazole
- Betamethasone valerate

Informacije o zdravilu

Ime zdravila:
OTOMAX EAR DROPS SUSPENSION

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
2640.00 international unit(s) / 1.00 millilitre(s)

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

Na voljo samo v English
1.07 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):

QS02CA90

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 v:

Sweden

Opis ovojnine zdravila:

Na voljo samo v Swedish
Na voljo samo v Swedish
Na voljo samo v Swedish
Na voljo samo v Swedish
Na voljo samo v Swedish
Na voljo samo v Swedish
Na voljo samo v Swedish
Na voljo samo v Swedish

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 French Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Intervet International B.V.

Datum dovoljenja za promet z zdravilom:
10/09/1999

Proizvodna mesta, odgovorna za sproščanje serij:
Trirx Segre

Pristojni organ:
Swedish Medical Products Agency

Številka dovoljenja za promet z zdravilom:
15571

Datum spremembe statusa dovoljenja za promet:
10/09/1999

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

Številka postopka:
FR/V/0110/001

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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Na voljo samo v Estonian English French Lithuanian Portuguese 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

Navodilo za uporabo

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

Povzetek glavnih značilnosti zdravila

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Označevanje

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Combined File of all Documents

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