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Mometamax Ultra 8.6 mg/ml + 2.6 mg/ml + 2.1 mg/ml - Ear drops, suspension

Ima dovoljenje za promet

- Gentamicin
- Posaconazole
- Mometasone furoate

Informacije o zdravilu

Ime zdravila:
Mometamax Ultra 8.6 mg/ml + 2.6 mg/ml + 2.1 mg/ml - 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
6880.00 international unit(s) / 1.00 millilitre(s)

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

Na voljo samo v English
1.68 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):

QS02CA91

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 samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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Na voljo v:
Austria , Belgium , Croatia , Cyprus , Czechia , France , Germany , Greece , Hungary , Ireland , Italy , Luxembourg , Netherlands , Poland , Romania , Slovakia , Slovenia , Spain

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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:
Intervet International B.V.

Datum dovoljenja za promet z zdravilom:
22/11/2022

Proizvodna mesta, odgovorna za sproščanje serij:
Vet Pharma Friesoythe GmbH

Pristojni organ:
European Commission

Številka dovoljenja za promet z zdravilom:
Ta podatek za to zdravilo ni na voljo.

Datum spremembe statusa dovoljenja za promet:
22/11/2022

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Combined File of all Documents

slovenščina (PDF)
Objavljeno na dan: 7/09/2026
Prenesi

ema-puar-mometamax-ultra-v-004987-par-en.pdf