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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)

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**Farmacevtska oblika:**

Kapljice za uho, suspenzija

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

QS02CA90

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**Pravni status za dobavo/izdajo zdravila:**

Zdravilo, ki se izdaja na veterinarski recept

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**Status dovoljenja za promet z zdravilom:**

Valid

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**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:**

Denmark

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**Opis ovojnine zdravila:**

Na voljo samo v French
Na voljo samo v French
Na voljo samo v French
Na voljo samo v French
Na voljo samo v French
Na voljo samo v French
Na voljo samo v French
Na voljo samo v French

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## 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:**
30/07/1999

**Proizvodna mesta, odgovorna za sproščanje serij:**
Trirx Segre

**Pristojni organ:**
Danish Medicines Agency

**Številka dovoljenja za promet z zdravilom:**
30828

**Datum spremembe statusa dovoljenja za promet:**
30/07/1999

**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
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

Povzetek glavnih značilnosti zdravila

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

Combined File of all Documents

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