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# Meloxidyl 0.5 mg/ml - Oral suspension

Ima dovoljenje za promet

- Meloxicam

## Informacije o zdravilu

**Ime zdravila:**
Meloxidyl 0.5 mg/ml - Oral suspension

**Učinkovina:**
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:**
Peroralna uporaba

## Podatki o zdravilu

**Učinkovina / Jakost:**
Na voljo samo v English
0.50 milligram(s) / 1.00 Steklenica/steklenička; Plastenka

**Farmacevtska oblika:**

Peroralna suspenzija

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

QM01AC06

**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 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 , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Portugal , Slovakia , Slovenia , Spain , Sweden ,

United Kingdom (Northern Ireland)

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**Opis ovojnine zdravila:**

Na voljo samo v English

Na voljo samo v English

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## Dodatne informacije

**Vrsta dovoljenja:**

Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

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**Pravna podlaga za izdajo dovoljenja za promet z zdravilom:**

Na voljo samo v English Italian Latvian Norwegian

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**Imetnik dovoljenja za promet z zdravilom:**

Ceva Sante Animale

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**Datum dovoljenja za promet z zdravilom:**

15/01/2007

**Proizvodna mesta, odgovorna za sproščanje serij:**
Ceva Sante Animale

**Pristojni organ:**
European Commission

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

**Datum spremembe statusa dovoljenja za promet:**
26/08/2010

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: 2/10/2026
Prenesi

ema-puar-meloxidyl-v-115-par-en.pdf