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Loxicom 1.5 mg/ml - Oral suspension (dogs)

Ima dovoljenje za promet

- Meloxicam

Informacije o zdravilu

Ime zdravila:
Loxicom 1.5 mg/ml - Oral suspension (dogs)

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
1.50 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 Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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Na voljo v:

Austria , Belgium , Cyprus , Czechia , Denmark , Estonia , Finland , Greece , Hungary , Ireland , Lithuania , Luxembourg , Malta , Netherlands , Poland , Portugal , Romania , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Opis ovojnine zdravila:

Na voljo samo v English

Na voljo samo v English

Na voljo samo v English

Na voljo samo v English

Na voljo samo v English

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

Imetnik dovoljenja za promet z zdravilom:

Norbrook Laboratories (Ireland) Limited

Datum dovoljenja za promet z zdravilom:
10/02/2009

Proizvodna mesta, odgovorna za sproščanje serij:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
10/02/2009

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: 14/03/2026
Prenesi

ema-puar-loxicom-v-141-var-x-0005-en.pdf

ema-puar-loxicom-v-141-var-x-0003-en.pdf

ema-puar-loxicom-v-141-var-x-0009-en.pdf

ema-puar-loxicom-v-141-par-en.pdf