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Trocoxil 30 mg - Chewable tablet

- Mavacoxib

Ima dovoljenje za promet

Informacije o zdravilu

Ime zdravila:
Trocoxil 30 mg - Chewable tablet
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
  30.00
  milligram(s)
  /
  1.00
  Tableta

Farmacevtska oblika:

- Žvečljiva tableta

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

- QM01AH92

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
- 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 Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
- Na voljo samo v Spanish Czech German Estonian Greek English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
- Na voljo samo v Spanish Czech German Estonian English French Italian Latvian 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 Spanish Czech German Estonian English French Italian Lithuanian Dutch Portuguese Romanian 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
- Na voljo samo v Spanish Czech German Estonian English French Croatian 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 Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Na voljo v:

- Austria
- Belgium
- Czechia
- Denmark
- Estonia
- Finland
- Germany
- Hungary
- Italy
- Latvia
- Lithuania
- Luxembourg
- Netherlands
- Poland
- Slovakia
- Spain
- Sweden
- United Kingdom (Northern Ireland)

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

Imetnik dovoljenja za promet z zdravilom:

- Zoetis Belgium

Datum dovoljenja za promet z zdravilom:

- 9/09/2008

Proizvodna mesta, odgovorna za sproščanje serij:

- Pfizer Italia S.r.l.

Pristojni organ:

- European Commission

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

- 9/09/2008

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

Dokumenti

Informacije o zdravilu

Combined File of all Documents

slovenščina (PDF)
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Prenesi
Drugi jeziki (24)
English (PDF)

Objavljeno na dan: 14/03/2026

Prenesi

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Prenesi

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Javno poro?ilo o oceni zdravila

ema-puar-trocoxil-v-132-par-en.pdf

English (PDF)
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