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Semintra 4 mg/ml - Oral solution

Ima dovoljenje za promet

- Telmisartan

Informacije o zdravilu

Ime zdravila:
Semintra 4 mg/ml - Oral solution

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
4.00 milligram(s) / 1.00 Steklenica/steklenička; Plastenka

Farmacevtska oblika:

Peroralna raztopina

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

QC09CA07

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 , Czechia , Denmark , Finland , France , Germany , Iceland , Luxembourg , Norway , Poland , Spain , Sweden

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Opis ovojnine zdravila:

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 Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:

Boehringer Ingelheim Vetmedica GmbH

Datum dovoljenja za promet z zdravilom:

13/02/2013

Proizvodna mesta, odgovorna za sproščanje serij:
Boehringer Ingelheim Vetmedica 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:
13/02/2013

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-semintra-v-2436-var-x-0008-en.pdf

ema-puar-v2436-semintra-initial-en.pdf

ema-puar-v2436-semintra-initial-en.pdf