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Simparica Trio 6 mg + 0.12 mg + 25 mg - Chewable tablet

Ima dovoljenje za promet

- Sarolaner
- Moxidectin
- Pyrantel embonate

Informacije o zdravilu

Ime zdravila:
Simparica Trio 6 mg + 0.12 mg + 25 mg - Chewable tablet

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:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
Presentation_strength:6 mg Reference:Hse Index:0
Na voljo samo v English
Presentation_strength:0.12 mg Reference:Ph. Eur. Index:1
Na voljo samo v English
Presentation_strength:25 mg Reference:Ph. Eur. Index:2

Farmacevtska oblika:
Žvečljiva tableta

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QP54AB52

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:
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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 , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Romania , Slovakia , Spain , United Kingdom (Northern Ireland)

Opis ovojnine zdravila:

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 Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Zoetis Belgium

Datum dovoljenja za promet z zdravilom:
17/09/2019

Proizvodna mesta, odgovorna za sproščanje serij:
Corden Pharma GmbH
Zoetis Belgium

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
17/09/2019

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: 13/03/2026
Prenesi

ema-puar-simparica-trio-v-4846-par-en.pdf

ema-puar-simparica-trio-v-004846-var-ii-0007-g-en.pdf

ema-puar-simparica-trio-v-4846-var-vra-0009-g-en.pdf

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