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Simparica 80 mg - Chewable tablet

Ima dovoljenje za promet

- Sarolaner

Informacije o zdravilu

Ime zdravila:
Simparica 80 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
Presentation_strength:80 mg Reference:In house Index:0

Farmacevtska oblika:

Žvečljiva tableta

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

QP53BE03

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 , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Romania , Slovakia , 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

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:
6/11/2015

Proizvodna mesta, odgovorna za sproščanje serij:
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:
6/11/2015

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-simparica-v-3991-par-en.pdf

ema-puar-simparicav-3991-var-ii-0006-en.pdf

ema-puar-simparica-mipet-easecto-var-ws-2217-en.pdf