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Profender 50 mg + 10 mg - Modified-release tablet

Ima dovoljenje za promet

- Emodepside
- Praziquantel
- Emodepside
- Praziquantel

Informacije o zdravilu

Ime zdravila:
Profender 50 mg + 10 mg - Modified-release tablet

Učinkovina:
Na voljo samo v English
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
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

Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
10.00 milligram(s) / 1.00 Tableta

Na voljo samo v English
50.00 milligram(s) / 1.00 Tableta

Na voljo samo v English
10.00 milligram(s) / 1.00 Tableta

Na voljo samo v English
50.00 milligram(s) / 1.00 Tableta

Farmacevtska oblika:

Tableta s prirejenim sproščanjem

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

QP52AA51

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 , France , Germany , Luxembourg , Netherlands , Poland , Portugal , Slovakia , Spain

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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
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

Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English

Imetnik dovoljenja za promet z zdravilom:
Vetoquinol S.A.

Datum dovoljenja za promet z zdravilom:
27/07/2005

Proizvodna mesta, odgovorna za sproščanje serij:
Vetoquinol S.A.
Vetoquinol S.A.
Vetoquinol S.A.
Vetoquinol S.A.
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Vetoquinol S.A.
Vetoquinol S.A.

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
25/08/2008

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: 12/11/2026
Prenesi

ema-puar-profender-v-097-par-en.pdf

ema-puar-profender-v-097-var-ii-0032-en.pdf