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Prazitel Plus Tablets for Dogs

Ima dovoljenje za promet

- Praziquantel
- Pyrantel embonate
- Febantel

Informacije o zdravilu

Ime zdravila:
Prazitel Plus 50+50+150 ΔΙΣΚΙΟ
Prazitel Plus Tablets for Dogs

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
50.00 milligram(s) / 1.00 Tableta

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

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

Farmacevtska oblika:

Tableta

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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Opis ovojnine zdravila:

Na voljo samo v English
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Na voljo samo v English
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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 Portuguese Norwegian

Imetnik dovoljenja za promet z zdravilom:
Chanelle Pharmaceuticals Manufacturing Limited

Datum dovoljenja za promet z zdravilom:
1/06/2010

Proizvodna mesta, odgovorna za sproščanje serij:
Chanelle Pharmaceuticals Manufacturing Limited

Pristojni organ:
National Organization For Medicines

Številka dovoljenja za promet z zdravilom:
61172/16-09-2015/K-0184101

Datum spremembe statusa dovoljenja za promet:
15/09/2015

Referenčna država članica:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Številka postopka:
IE/V/0241/001

Zadevne države članice:

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 Latvian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Povzetek glavnih značilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenščina). Spodaj ga najdete v drugem jeziku.