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Drontal Dog Tasty Bone XL 525/504/175 mg tablets

Ima dovoljenje za promet

- Febantel
- Pyrantel embonate
- Praziquantel

Informacije o zdravilu

Ime zdravila:
Drontal Dog Tasty Bone XL 525/504/175 mg tablets

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
525.00 milligram(s) / 1.00 Tableta

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

Na voljo samo v English
175.00 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

Na voljo v:

Germany

Opis ovojnine zdravila:

Na voljo samo v Swedish
Na voljo samo v Swedish
Na voljo samo v Swedish
Na voljo samo v Swedish
Na voljo samo v Swedish

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 Italian Latvian Norwegian

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

Datum dovoljenja za promet z zdravilom:
1/09/2017

Proizvodna mesta, odgovorna za sproščanje serij:
KVP Pharma+Veterinaer Produkte GmbH
Europeenne De Pharmacotechnie Europhartech

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
402370.00.00

Datum spremembe statusa dovoljenja za promet:
11/10/2022

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

Številka postopka:
IE/V/0335/002

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 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

Combined File of all Documents

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