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PestiGon Combo 134 mg / 120.6 mg spot-on solution for medium dogs

Ima dovoljenje za promet

- Fipronil
- (S)-Methoprene

Informacije o zdravilu

Ime zdravila:
PestiGon Combo 134 mg / 120.6 mg spot-on solution for medium dogs

Učinkovina:
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:
Nanos na kožo

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
134.00 milligram(s) / 1.00 Pipeta

Na voljo samo v English
120.60 milligram(s) / 1.00 Pipeta

Farmacevtska oblika:

Kožni nanos, raztopina

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

QP53AX65

Pravni status za dobavo/izdajo zdravila:

Zdravilo, ki se izdaja brez veterinarskega recepta

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
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
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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 Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Norbrook Laboratories (Ireland) Limited

Datum dovoljenja za promet z zdravilom:
19/05/2017

Proizvodna mesta, odgovorna za sproščanje serij:
Norbrook Manufacturing Limited
Norbrook Laboratories Limited

Pristojni organ:
Directorate Of Veterinary Medicinal Products

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

Datum spremembe statusa dovoljenja za promet:
19/05/2017

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

Številka postopka:
IE/V/0363/003

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 German Estonian English French Italian Dutch Portuguese Slovak Finnish Swedish Icelandic Norwegian
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Na voljo samo v Estonian English French Lithuanian Portuguese 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). Lahko ga najdete spodaj v drugem jeziku.