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FRONTLINE COMBO 268.00 MG / 241.20 MG SPOT-ON SOLUTION FOR DOG L

Ima dovoljenje za promet

- Fipronil
- Methoprene

Informacije o zdravilu

Ime zdravila:
FRONTLINE COMBO 268.00 MG / 241.20 MG SPOT-ON SOLUTION FOR DOG L

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:
Dermalna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

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

Na voljo samo v English
241.20 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 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:

Norway

Opis ovojnine zdravila:

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 Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Boehringer Ingelheim Animal Health Denmark A/S

Datum dovoljenja za promet z zdravilom:
10/02/2009

Proizvodna mesta, odgovorna za sproščanje serij:
Boehringer Ingelheim Animal Health France

Pristojni organ:
Norwegian Medical Products Agency

Številka dovoljenja za promet z zdravilom:
03-1993

Datum spremembe statusa dovoljenja za promet:
10/02/2009

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

Številka postopka:
FR/V/0139/003

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 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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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.

Navodilo za uporabo

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Combined File of all Documents

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