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DYNACAN 50 mg/60mg SPOT-ON SOLUTION CATS AND FERRETS

Ima dovoljenje za promet

- Fipronil
- Methoprene

Informacije o zdravilu

Ime zdravila:
DYNACAN 50 mg/60mg SPOT-ON SOLUTION CATS AND FERRETS

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
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
50.00 milligram(s) / 1.00 Pipeta

Na voljo samo v English
60.00 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

Dodatne informacije

Vrsta dovoljenja:

Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

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Imetnik dovoljenja za promet z zdravilom:
Calier Portugal Medicamentos E Produtos Veterinarios S.A.

Datum dovoljenja za promet z zdravilom:
17/01/2020

Proizvodna mesta, odgovorna za sproščanje serij:
Laboratorios Calier S.A.

Pristojni organ:
Directorate General For Food And Veterinary

Številka dovoljenja za promet z zdravilom:
1330/01/20DFVPT

Datum spremembe statusa dovoljenja za promet:
19/10/2022

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

Številka postopka:
PT/V/0130/001

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 Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Lithuanian Dutch Portuguese Romanian Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet