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Vectra 3D 436 mg dinotefuran, 38.7 mg pyriproxyfen, 3175 mg permethrin - Spot-on solution

Ima dovoljenje za promet

- Permethrin
- Pyriproxyfen
- Dinotefuran

Informacije o zdravilu

Ime zdravila:
Vectra 3D 436 mg dinotefuran, 38.7 mg pyriproxyfen, 3175 mg permethrin - Spot-on solution

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:
Nanos na kožo

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
Presentation_strength:3175 mg Reference:Hse Index:0
Na voljo samo v English
Presentation_strength:38.7 mg Reference:Hse Index:1
Na voljo samo v English
Presentation_strength:436 mg Reference:Hse Index:2

Farmacevtska oblika:
Kožni nanos, raztopina

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QP53AC54

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:
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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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Na voljo v:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Luxembourg , Malta , Netherlands , Poland , Portugal , Romania , Slovakia , Spain , Sweden ,

United Kingdom (Northern Ireland)

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
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

Imetnik dovoljenja za promet z zdravilom:
Ceva Sante Animale

Datum dovoljenja za promet z zdravilom:
4/12/2013

Proizvodna mesta, odgovorna za sproščanje serij:
Ceva Sante Animale
Ab7 Sante

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
4/12/2013

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

Dokumenti

Combined File of all Documents

slovenščina (PDF)
Objavljeno na dan: 30/10/2026
Prenesi

ema-puar-vectra-3d-v-2555-par-en.pdf

ema-puar-vectra-3d-v-002555-referral-a82-0023-en.pdf