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NexGard Combo 3.6 mg + 1.2 mg + 24.9 mg - Spot-on solution

Ima dovoljenje za promet

- Esafoxolaner
- Eprinomectin
- Praziquantel

Informacije o zdravilu

Ime zdravila:
NexGard Combo 3.6 mg + 1.2 mg + 24.9 mg - 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:3.6 mg Reference:Hse Index:0
Na voljo samo v English
Presentation_strength:1.2 mg Reference:Hse Index:1
Na voljo samo v English
Presentation_strength:24.9 mg Reference:Ph.Eur. Index:2

Farmacevtska oblika:
Kožni nanos, raztopina

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QP54AA54

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:
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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 , Czechia , Denmark , Estonia , Finland , France , Latvia , Lithuania , Luxembourg , Norway , Poland , Slovakia , Spain , Sweden

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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

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:
Boehringer Ingelheim Vetmedica GmbH

Datum dovoljenja za promet z zdravilom:
6/01/2021

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

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
6/01/2021

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: 13/03/2026

Prenesi

ema-puar-v5094-nexgard-combo-vra0007g-en.pdf

ema-puar-nexgard-combo-v-5094-par-en.pdf

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