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Advocate 250 mg + 62.5 mg - Spot-on solution

Ima dovoljenje za promet

- Imidacloprid
- Moxidectin

Informacije o zdravilu

Ime zdravila:
Advocate 250 mg + 62.5 mg - Spot-on solution

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
Presentation_strength:250 mg Reference:Hse Index:0

Na voljo samo v English
Presentation_strength:62.5 mg Reference:In-house Index:1

Farmacevtska oblika:
Kožni nanos, raztopina

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QP54AB52

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
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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 , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Opis ovojnine zdravila:

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
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

Imetnik dovoljenja za promet z zdravilom:
Elanco Animal Health GmbH

Datum dovoljenja za promet z zdravilom:
2/04/2003

Proizvodna mesta, odgovorna za sproščanje serij:
KVP Pharma+Veterinaer Produkte GmbH

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
2/04/2003

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: 14/03/2026

Prenesi

ema-puar-advocate-v-076-var-ii-0041-g-en.pdf

ema-puar-advocate-v-076-var-ii-0026-g-en.pdf

ema-puar-advocate-v-076-par-en.pdf

ema-puar-advocate-v-076-var-ii-0039-g-en.pdf

ema-puar-advocate-v-076-var-ii-0022-en.pdf

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