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VarroMed 44 mg/ml + 5 mg/ml - bee-hive dispersion

Ima dovoljenje za promet

- Oxalic acid dihydrate
- Formic acid

Informacije o zdravilu

Ime zdravila:
VarroMed 44 mg/ml + 5 mg/ml - bee-hive dispersion

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:
Dajanje v čebelji panj

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
43.75 milligram(s) / 1.00 Steklenica/steklenička; Plastenka

Na voljo samo v English
5.09 milligram(s) / 1.00 Steklenica/steklenička; Plastenka

Farmacevtska oblika:

Disperzija za uporabo v čebeljem panju

Karenca glede na pot uporabe:

Dajanje v čebelji panj:

- **Honey bee**
  - Honey. 0 day Zero days

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

QP53AG30

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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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 , France , Germany , Greece , Hungary , Ireland , Italy , Lithuania , Luxembourg , Netherlands , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , United Kingdom (Northern Ireland)

Opis ovojnine zdravila:

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:
BeeVital GmbH

Datum dovoljenja za promet z zdravilom:
2/02/2017

Proizvodna mesta, odgovorna za sproščanje serij:
Lichtenheldt GmbH Pharmazeutische Fabrik
Labor LS SE & Co. KG

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
2/02/2017

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: 15/10/2026
Prenesi

ema-puar-varromed-v-2723-par-en.pdf