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Thymovar

Ima dovoljenje za promet

- Thymol

Informacije o zdravilu

Ime zdravila:
Thymovar

Učinkovina:
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
15.00 gram(s) / 1.00 Trak

Farmacevtska oblika:
Trak za uporabo v čebeljem panju

Karenca glede na pot uporabe:

Dajanje v čebelji panj:

- **Honey bee**
  - Honey. 0 day

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

QP53AX22

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 v:

Netherlands

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

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Pravna podlaga za izdajo dovoljenja za promet z zdravilom:

Na voljo samo v English Italian Latvian Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:

Andermatt BioVet GmbH

Datum dovoljenja za promet z zdravilom:
27/09/2006

Proizvodna mesta, odgovorna za sproščanje serij:
Andermatt BioVet GmbH

Pristojni organ:
Medicines Evaluation Board

Številka dovoljenja za promet z zdravilom:
REG NL 10330

Datum spremembe statusa dovoljenja za promet:
25/01/2022

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

Številka postopka:
NL/V/0120/001

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

Dokumenti

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.