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APIVAR 500 MG AMITRAZ BEE-HIVE STRIPS FOR HONEY BEES

Ima dovoljenje za promet

- Amitraz

Informacije o zdravilu

Ime zdravila:
APIVAR 500 MG AMITRAZ BEE-HIVE STRIPS FOR HONEY BEES

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
0.50 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

Do not use during honey flow. Do not extract honey from the brood chamber. Do not harvest honey when the treatment is in place. Brood combs should be replaced with new foundation at least every three years. Do not recycle brood frames as honey frames.

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QP53AD01

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

Opis ovojnine zdravila:
Na voljo samo v Swedish
Na voljo samo v Swedish
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:
Veto-Pharma

Datum dovoljenja za promet z zdravilom:
24/11/2017

Proizvodna mesta, odgovorna za sproščanje serij:
Veto-Pharma

Pristojni organ:
Swedish Medical Products Agency

Številka dovoljenja za promet z zdravilom:
56256

Datum spremembe statusa dovoljenja za promet:
24/11/2017

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

Številka postopka:
FR/V/0319/001

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
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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

Navodilo za uporabo

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

Označevanje

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

Povzetek glavnih značilnosti zdravila

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Combined File of all Documents

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