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Apilife Var bee-hive strip for honey bees

Ima dovoljenje za promet

- Thymol
- EUCALYPTUS OIL
- Camphor, racemic
- Levomenthol

Informacije o zdravilu

Ime zdravila:
Apilife Var bee-hive strip for honey bees

Učinkovina:
Na voljo samo v English
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:
Dajanje v čebelji panj

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
8.00 gram(s) / 1.00 Trak

Na voljo samo v English
1.72 gram(s) / 1.00 Trak

Na voljo samo v English
0.39 gram(s) / 1.00 Trak

Na voljo samo v English
0.39 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 in order to avoid to adulterate the taste of the honey

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

QP53AX30

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:
Germany

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

Imetnik dovoljenja za promet z zdravilom:
Chemicals Laif S.p.A.

Datum dovoljenja za promet z zdravilom:
18/08/2009

Proizvodna mesta, odgovorna za sproščanje serij:
Chemicals Laif S.p.A.

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
401267.00.00

Datum spremembe statusa dovoljenja za promet:
5/11/2014

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

Številka postopka:
IT/V/0123/001

Zadevne države članice:

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