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AFILARIA SR 3.4 mg/ml powder and solvent for suspension for injection for dogs

Ima dovoljenje za promet

- Moxidectin

Informacije o zdravilu

Ime zdravila:
AFILARIA SR 3.4 mg/ml powder and solvent for suspension for injection for dogs

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:
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
100.00 milligram(s)/gram / 1.00 gram(s)

Farmacevtska oblika:
Prašek in vehikel za suspenzijo za injiciranje

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QP54AB02

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

Na voljo v:
Austria

Opis ovojnine zdravila:
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 Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Fatro S.p.A.

Datum dovoljenja za promet z zdravilom:

21/06/2022

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

Pristojni organ:
Austrian Agency For Health And Food Safety

Številka dovoljenja za promet z zdravilom:
841264

Datum spremembe statusa dovoljenja za promet:
21/06/2022

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

Številka postopka:
ES/V/0315/001

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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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.

Povzetek glavnih značilnosti zdravila

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

Označevanje

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Combined File of all Documents

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