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Amcofen 4 mg/10 mg chewable tablets for small cats and kittens weighing at least 0.5 kg

Ima dovoljenje za promet

- Milbemycin oxime
- Praziquantel

Informacije o zdravilu

Ime zdravila:
Amcofen 4 mg/10 mg chewable tablets for small cats and kittens weighing at least 0.5 kg

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:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
4.00 milligram(s) / 1.00 Tableta

Na voljo samo v English
10.00 milligram(s) / 1.00 Tableta

Farmacevtska oblika:

Žvečljiva tableta

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

QP54AB51

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

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Na voljo v:

Germany

Opis ovojnine zdravila:

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

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Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
KRKA tovarna zdravil d.d. Novo mesto

Datum dovoljenja za promet z zdravilom:
13/04/2019

Proizvodna mesta, odgovorna za sproščanje serij:
TAD Pharma GmbH
Krka-Farma d.o.o.
KRKA tovarna zdravil d.d. Novo mesto

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
402636.00.00

Datum spremembe statusa dovoljenja za promet:
13/04/2019

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

Številka postopka:
IE/V/0525/001

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

Combined File of all Documents

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