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MILBEMAX 2.5 MG/25 MG CHEWABLE TABLETS FOR SMALL DOGS AND PUPPIES

Ima dovoljenje za promet

- Milbemycin oxime
- Praziquantel

Informacije o zdravilu

Ime zdravila:
MILBEMAX 2.5 MG/25 MG CHEWABLE TABLETS FOR SMALL DOGS AND PUPPIES

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
2.50 milligram(s) / 1.00 Tableta

Na voljo samo v English
25.00 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:
Na voljo samo v English French Italian Latvian Lithuanian Portuguese Finnish Swedish Norwegian

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:
France

Opis ovojnine zdravila:
Na voljo samo v French
Na voljo samo v French
Na voljo samo v French
Na voljo samo v French
Na voljo samo v French
Na voljo samo v French

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 Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Elanco GmbH

Datum dovoljenja za promet z zdravilom:
30/01/2009

Proizvodna mesta, odgovorna za sproščanje serij:
Elanco France S.A.S.

Pristojni organ:
French Agency For Food, Environmental And Occupational Health & Safety

Številka dovoljenja za promet z zdravilom:
FR/V/4735370 3/2009

Datum spremembe statusa dovoljenja za promet:
30/01/2014

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

Številka postopka:
FR/V/0135/005

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 Greek English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Dokumenti

Povzetek glavnih značilnosti zdravila

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Package Leaflet and Labelling

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Combined File of all Documents

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