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Source URL: https://medicines.health.europa.eu/veterinary/sl/600000080421
Mektix 12.5 mg/125 mg film-coated tablets for dogs weighing at least 5 kg

- Milbemycin oxime
- Praziquantel

Ima dovoljenje za promet

Informacije o zdravilu

Ime zdravila:
Mektix 12.5 mg/125 mg film-coated tablets for dogs weighing at least 5 kg
U?inkovina:

- 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

U?inkovina / Jakost:

- Na voljo samo v English
  12.50
  milligram(s)
  /
  1.00
  Tableta
- Na voljo samo v English
  125.00
  milligram(s)
  /
  1.00
  Tableta

Farmacevtska oblika:

- Filmsko obložena tableta

Anatomsko-terapevtsko-kemi?na 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:

- Sweden

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

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:

- 5/01/2022

Proizvodna mesta, odgovorna za sproš?anje serij:

- KRKA tovarna zdravil d.d. Novo mesto
- Krka-Farma d.o.o.

Pristojni organ:

- Swedish Medical Products Agency

Številka dovoljenja za promet z zdravilom:

- 61019

Datum spremembe statusa dovoljenja za promet:

- 5/01/2022

Referen?na država ?lanica:

- Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Številka postopka:

- IE/V/0526/004

Zadevne države ?lanice:

- 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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- Na voljo samo v Spanish Czech German Estonian English French Italian Lithuanian Dutch Portuguese Romanian Slovak Swedish Icelandic Norwegian
- Na voljo samo v Spanish Czech German Estonian English French Croatian 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 Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Za poro?ila o domnevnih neželenih u?inkih zdravil za uporabo v veterinaski medicini obiš?ite stran: www.adrreports.eu/vet

Dokumenti

Informacije o zdravilu

Povzetek glavnih zna?ilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenš?ina). Lahko ga najdete spodaj v drugem jeziku.
Drugi jeziki (2)
Swedish (PDF)
Objavljeno na dan: 8/09/2023
Updated on: 14/03/2026
Prenesi
English (PDF)
Objavljeno na dan: 30/06/2024
Updated on: 13/03/2026
Prenesi

Navodilo za uporabo

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Updated on: 14/03/2026
Prenesi

Ozna?evanje

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Drugi jeziki (1)
Swedish (PDF)
Objavljeno na dan: 8/09/2023
Updated on: 14/03/2026
Prenesi

Javno poro?ilo o oceni zdravila

ie-puar-mr-iev0526004-mektix-125-mg125-mg-film-coated-tablets-for-dogs-w-en.pdf

English (PDF)
Objavljeno na dan: 26/08/2024
Updated on: 14/03/2026
Prenesi