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VETMULIN 125 mg/ml Solution for use in drinking water for pigs and chickens

Ima dovoljenje za promet

- Tiamulin hydrogen fumarate
- Tiamulin hydrogen fumarate

Informacije o zdravilu

Ime zdravila:
VETMULIN 125 mg/ml Solution for use in drinking water for pigs and chickens

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 Italian Latvian Lithuanian Hungarian Romanian Finnish Swedish Icelandic
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 vodo za pitje

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
125.00 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
125.00 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:

Raztopina za dajanje v vodo za pitje

Karenca glede na pot uporabe:

Dajanje v vodo za pitje:

- **Poultry**
  - Meat and offal. 2 day
  - Meat and offal. 2 day
  - Eggs. 0 day
  - Eggs. 0 day
- **Pig**
  - Meat and offal. 2 day

8,8 mg tiamulinhydrogenfumarat/kg kropsvægt svarende til 7 ml af produktet/100 kg kropsvægt

- Meat and offal. 2 day

8,8 mg tiamulinhydrogenfumarat/kg kropsvægt svarende til 7 ml af produktet/100 kg kropsvægt

- Meat and offal. 4 day

20 mg tiamulinhydrogenfumarat/kg kropsvægt svarende til 16 ml af produktet/100 kg kropsvægt

- Meat and offal. 4 day

20 mg tiamulinhydrogenfumarat/kg kropsvægt svarende til 16 ml af produktet/100 kg kropsvægt

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QJ01XQ01

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:
Denmark

Opis ovojnine zdravila:
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:
HuVepharma

Datum dovoljenja za promet z zdravilom:

19/02/2019

Proizvodna mesta, odgovorna za sproščanje serij:
Biovet AD

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
60221

Datum spremembe statusa dovoljenja za promet:
19/02/2019

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

Številka postopka:
DK/V/0122/001

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 Spanish Czech German Estonian English French 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

Povzetek glavnih značilnosti zdravila

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

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