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GABBROVET MULTI 140 MG/ML SOLUTION FOR USE IN DRINKING WATER/ MILK FOR PRE-RUMINANT CATTLE AND PIGS

Ima dovoljenje za promet

- Paromomycin sulfate

Informacije o zdravilu

Ime zdravila:
GABBROVET MULTI 140 MG/ML SOLUTION FOR USE IN DRINKING WATER/ MILK FOR PRE-RUMINANT CATTLE AND PIGS

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
tele
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
200.00 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Raztopina za dajanje v vodo za pitje/mleko

Karenca glede na pot uporabe:
Peroralna uporaba:

- **tele**
  - Meat and offal. 110 day

Cryptosporidiosis: Dosage: 150 mg/kg/day for 5 days.

  - Meat and offal. 20 day

Colibacillosis: Dosage: 25-50 mg/kg/day for 3 to 5 days.

- **Pig**
  - Meat and offal. 3 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QA07AA06

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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Opis ovojnine zdravila:
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
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:
Ceva Sante Animale

Datum dovoljenja za promet z zdravilom:
24/08/2022

Proizvodna mesta, odgovorna za sproščanje serij:
Ceva Sante Animale

Pristojni organ:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Številka dovoljenja za promet z zdravilom:
DC/V/0614/002

Datum spremembe statusa dovoljenja za promet:
24/08/2022

Referenčna država članica:

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

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Številka postopka:
FR/V/0429/001

Zadevne države članice:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Povzetek glavnih značilnosti zdravila

Package Leaflet and Labelling

Combined File of all Documents

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

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