File downloaded on 2026-11-04
Source URL: https://medicines.health.europa.eu/veterinary/en/600000031094

GABBROVET 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 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:
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Norwegian
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/mleko

Dajanje v vodo za pitje

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:

Dajanje v vodo za pitje/mleko:

- **Cattle (pre-ruminant)**
  - Meat and offal. 20 day

Dajanje v vodo za pitje:

- **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 German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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
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:
14/09/2018

Proizvodna mesta, odgovorna za sproščanje serij:
Ceva Sante Animale

Pristojni organ:
Icelandic Medicines Agency

Številka dovoljenja za promet z zdravilom:
IS/2/18/009/01

Datum spremembe statusa dovoljenja za promet:
14/09/2018

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

Številka postopka:
FR/V/0317/001

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

---

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.

eu-puar-frv0317001-mr-rpe364-en.pdf