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NEOMAY 500 000 IU/G POWDER FOR USE IN DRINKING WATER/MILK REPLACER

Ima dovoljenje za promet

- Neomycin

Informacije o zdravilu

Ime zdravila:
NEOMAY 500 000 IU/G POWDER FOR USE IN DRINKING WATER/MILK REPLACER

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
govedo
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English Italian Latvian Lithuanian Hungarian Romanian Finnish Swedish Icelandic

Pot uporabe:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
500000.00 international unit(s) / 1.00 gram(s)

Farmacevtska oblika:

Prašek za dajanje v vodo za pitje/mleko

Karenca glede na pot uporabe:

Peroralna uporaba:

- **govedo**
  - Meat and offal. 14 day
- **Pig**
  - Meat and offal. 3 day
- **Poultry**
  - Meat and offal. 14 day
  - Eggs. 0 day

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

QA07AA01

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

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:
Laboratorios Maymo S.A.U.

Datum dovoljenja za promet z zdravilom:
1/12/2015

Proizvodna mesta, odgovorna za sproščanje serij:
Laboratorios Maymo S.A.U.

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
54739

Datum spremembe statusa dovoljenja za promet:
1/12/2015

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

Številka postopka:
FR/V/0282/001

Zadevne države članice:

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 Latvian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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 Estonian English French Lithuanian Portuguese 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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