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Laksans 500 g peroralni prašek

Ni odobreno

- Magnesium sulfate heptahydrate

Informacije o zdravilu

Ime zdravila:
Laksans 500 g peroralni prašek

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 French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
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 French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Pot uporabe:
Dajanje v vodo za pitje/mleko
Dajanje v krmno mešanico

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
500.00 gram(s) / 1.00 Vreča

Farmacevtska oblika:

Peroralni prašek

Karenca glede na pot uporabe:

Dajanje v vodo za pitje/mleko:

- **govedo**
  - Meat and offal, milk. 0 day Meso in organi: nič dni. Mleko: nič dni.
- **Horse**
  - Meat and offal, milk. 0 day
- **Sheep**
  - Meat and offal, milk. 0 day
- **Goat**
  - Meat and offal, milk. 0 day
- **Pig**
  - Meat and offal. 0 day

Dajanje v krmno mešanico:

- **govedo**
  - Meat and offal, milk. 0 day
-

Horse
- Meat and offal, milk. 0 day

•

Sheep
- Meat and offal, milk. 0 day

•

Goat
- Meat and offal, milk. 0 day

•

Pig
- Meat and offal. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QA06AD04

Pravni status za dobavo/izdajo zdravila:
Zdravilo, ki se izdaja brez veterinarskega recepta

Status dovoljenja za promet z zdravilom:
Surrendered

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:
Laksans peroralni prašek je pakiran v PE vrečke po 500 g. Vsaka vrečka je v posamezni kartonski škatli.

Dodatne informacije

Vrsta dovoljenja:
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

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Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English French Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
GENERA SI podjetje za zastopanje in trgovino d.o.o.

Datum dovoljenja za promet z zdravilom:
29/11/2021

Proizvodna mesta, odgovorna za sproščanje serij:
Genera d.d.

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

Številka dovoljenja za promet z zdravilom:
NP/V/0184/001

Datum spremembe statusa dovoljenja za promet:
25/11/2022

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