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RABISIN, suspensie injectabilă

Ima dovoljenje za promet

- Rabies virus, strain G52, Inactivated

Informacije o zdravilu

Ime zdravila:
RABISIN, suspensie injectabilă

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
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
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

Pot uporabe:

Subkutana uporaba

intramuskularna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
2.09 log10 optical density50 / 1.00 millilitre(s)

Farmacevtska oblika:

Suspenzija za injiciranje

Karenca glede na pot uporabe:

Subkutana uporaba:

- **govedo**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day

intramuskularna uporaba:

- **govedo**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI07AA02

Pravni status za dobavo/izdajo zdravila:
Ta podatek za to zdravilo ni na voljo.

Status dovoljenja za promet z zdravilom:
Valid

Ima dovoljenje za promet v:
Na voljo samo v Spanish Czech German Estonian English French Italian Lithuanian Dutch Portuguese Romanian Slovak Swedish Icelandic Norwegian

Na voljo v:
Romania

Opis ovojnine zdravila:
Na voljo samo v Romanian
Na voljo samo v Romanian
Na voljo samo v Romanian
Na voljo samo v Romanian

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

Imetnik dovoljenja za promet z zdravilom:
Boehringer Ingelheim Animal Health France

Datum dovoljenja za promet z zdravilom:
30/08/1999

Proizvodna mesta, odgovorna za sproščanje serij:

Boehringer Ingelheim Animal Health France

Pristojni organ:
Institute For Control Of Biological Products And Veterinary Medicines

Številka dovoljenja za promet z zdravilom:
150136

Datum spremembe statusa dovoljenja za promet:
22/06/2025

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.