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Rabisin Injektionssuspension für Hunde, Katzen, Marderartige, Pferde, Rinder und Schafe

Ima dovoljenje za promet

- Rabies virus, strain G52, Inactivated

Informacije o zdravilu

Ime zdravila:
Rabisin Injektionssuspension für Hunde, Katzen, Marderartige, Pferde, Rinder und Schafe

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
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 Danish German Estonian English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Norwegian

Pot uporabe:
intramuskularna uporaba
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
1.00 international unit(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Suspenzija za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

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

Subkutana uporaba:

-

govedo

- Milk. 0 day
- Meat and offal. 0 day

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Sheep

- Meat and offal. 0 day
- Milk. 0 day

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

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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Na voljo v:
Germany

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

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 Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Boehringer Ingelheim Vetmedica GmbH

Datum dovoljenja za promet z zdravilom:
24/11/2005

Proizvodna mesta, odgovorna za sproščanje serij:
Boehringer Ingelheim Animal Health France

Pristojni organ:
Paul-Ehrlich-Institut

Številka dovoljenja za promet z zdravilom:
94a/84

Datum spremembe statusa dovoljenja za promet:
20/07/2010

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.