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RABIROM I

Ni odobreno

- Rabies virus, strain CVS-11, Inactivated

Informacije o zdravilu

Ime zdravila:
RABIROM I

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
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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
2.00 international unit(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Suspenzija za injiciranje

Karenca glede na pot uporabe:
Subkutana uporaba:

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

intramuskularna uporaba:

-

Pig

- Meat and offal. 0 day

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

QI07XA

Pravni status za dobavo/izdajo zdravila:

Ta podatek za to zdravilo ni na voljo.

Status dovoljenja za promet z zdravilom:

Surrendered

Ima dovoljenje za promet v:

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

Opis ovojnine zdravila:

Na voljo samo v Romanian
Na voljo samo v Romanian
Na voljo samo v Romanian
Na voljo samo v Romanian
Na voljo samo v Romanian
Na voljo samo v Romanian
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:
Romvac Company S.A.

Datum dovoljenja za promet z zdravilom:
13/08/2019

Proizvodna mesta, odgovorna za sproščanje serij:
Romvac Company S.A.

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

Številka dovoljenja za promet z zdravilom:
190215

Datum spremembe statusa dovoljenja za promet:
21/07/2024

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.