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# RABISIN SUSPENSION INJECTABLE

Ima dovoljenje za promet

- Rabies virus, strain G52, Inactivated

## Informacije o zdravilu

**Ime zdravila:**
RABISIN SUSPENSION INJECTABLE

**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
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:**

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:**

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

**Subkutana uporaba:**

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

**Na voljo v:**
France

**Opis ovojnine zdravila:**
Na voljo samo v French
Na voljo samo v French
Na voljo samo v French

## 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 French Italian Latvian Norwegian

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

**Datum dovoljenja za promet z zdravilom:**
7/01/1981

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

**Pristojni organ:**
French Agency For Food, Environmental And Occupational Health & Safety

**Številka dovoljenja za promet z zdravilom:**
FR/V/5411888 9/1981

**Datum spremembe statusa dovoljenja za promet:**
7/01/2011

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.

Package Leaflet and Labelling

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.