Rifen 100 mg/ml - Injektionslösung für Pferde, Rinder und Schweine

Pooblaščeno

- Ketoprofen

Product identification

Ime zdravila:
Rifen 100 mg/ml - Injektionslösung für Pferde, Rinder und Schweine
Rifen 100 mg/ml oplossing voor injectie voor paarden, runderen en varkens

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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 Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Swedish Icelandic Norwegian

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Pot uporabe:
intramuskularna uporaba
Intravenska uporaba

Product details

Učinkovina / Jakost:

Na voljo samo v English
100.00 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:

Raztopina za injiciranje

Withdrawal period by route of administration:

intramuskularna uporaba:

- **govedo**

  - Meat and offal. 3 day

  - Milk. 0 hour

- **Pig**

  - Meat and offal. 4 day

Intravenska uporaba:

- **govedo**

  - Meat and offal. 1 day

  - Milk. 0 hour

- **Horse**

  - Meat and offal. 1 day

Anatomsko-terapevtsko-kemična veterinarska oznaka (ATCvet):

QM01AE03

Pravni status za dobavo / izdajo zdravila:

Zdravilo, ki se izdaja na veterinarski recept

Status dovoljenja:

Valid

Authorised in:

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

Opis ovojnine:

Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

Additional information

Entitlement type:

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 Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:

Vetviva Richter GmbH

Marketing authorisation date:

21/12/2012

Proizvodna mesta, odgovorna za sproščanje serij:

Vetviva Richter GmbH

Pristojni organ:

Medicines Evaluation Board

Številka dovoljenja :

REG NL 105079

Datum spremembe statusa dovoljenja:

18/01/2022

Referenčna država članica:

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

Številka postopka:

Zadevne države članice:

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

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Na voljo samo v Estonian English French Swedish Icelandic Norwegian

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

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

Source URL: *https://medicines.health.europa.eu/veterinary/600000015294*