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# Rivalgin, 500mg/ml, Solution for injection

Ima dovoljenje za promet

- Metamizole sodium monohydrate

## Informacije o zdravilu

**Ime zdravila:**
Rivalgin, 500mg/ml, Solution for injection

**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

**Pot uporabe:**
intramuskularna uporaba
Intravenska uporaba

# Podatki o zdravilu

**Učinkovina / Jakost:**

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

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**Farmacevtska oblika:**

Raztopina za injiciranje

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**Karenca glede na pot uporabe:**

**intramuskularna uporaba:**

- **govedo**
  - Milk. 48 hour
  - Meat and offal. 12 day
- **Pig**
  - Meat and offal. 12 day

**Intravenska uporaba:**

- **Horse**
  - Milk. no withdrawal period

Not authorised for use in mares producing milk for human consumption.,

  - Meat and offal. 5 day

- **govedo**
  - Milk. 48 hour
  - Meat and offal. 12 day
- **Pig**
  - Meat and offal. 12 day

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**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QN02BB02

**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:**
Denmark

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

## 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 Italian Latvian Norwegian

**Imetnik dovoljenja za promet z zdravilom:**
Vetviva Richter GmbH

**Datum dovoljenja za promet z zdravilom:**
14/08/2019

**Proizvodna mesta, odgovorna za sproščanje serij:**
Vetviva Richter GmbH

**Pristojni organ:**

Danish Medicines Agency

**Številka dovoljenja za promet z zdravilom:**
61671

**Datum spremembe statusa dovoljenja za promet:**
14/08/2019

**Referenčna država članica:**
Na voljo samo v Spanish Czech German Estonian English French Italian Latvian Dutch Portuguese Slovak Swedish Icelandic Norwegian

**Številka postopka:**
CZ/V/0152/001

**Zadevne države članice:**
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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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.

Označevanje

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

Navodilo za uporabo

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

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