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Quiflor 100 mg/ml solution for injection for cattle and pigs (sows)

Ima dovoljenje za promet

- Marbofloxacin

Informacije o zdravilu

Ime zdravila:
Quiflor 100 mg/ml solution for injection for cattle and pigs (sows)

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
govedo
Na voljo samo v Bulgarian Spanish Danish German Estonian English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Pot uporabe:
Intravenska uporaba
Subkutana uporaba
intramuskularna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

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

Farmacevtska oblika:

Raztopina za injiciranje

Karenca glede na pot uporabe:

Intravenska uporaba:

- **govedo**
  - Milk. 36 hour
  - Meat and offal. 6 day

Subkutana uporaba:

- **govedo**
  - Milk. 36 hour
  - Meat and offal. 6 day

intramuskularna uporaba:

- **govedo**
  - Milk. 36 hour
  - Meat and offal. 6 day
- **Pig (sow for reproduction)**
  - Meat and offal. 4 day

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

QJ01MA93

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

Opis ovojnine zdravila:
Na voljo samo v English
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:
KRKA tovarna zdravil d.d. Novo mesto

Datum dovoljenja za promet z zdravilom:
19/05/2011

Proizvodna mesta, odgovorna za sproščanje serij:
KRKA tovarna zdravil d.d. Novo mesto
TAD Pharma GmbH

Pristojni organ:
Directorate General For Food And Veterinary

Številka dovoljenja za promet z zdravilom:
349/01/11DFVPT

Datum spremembe statusa dovoljenja za promet:
30/06/2025

Referenčna država članica:

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

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Številka postopka:
DE/V/0302/001

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
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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.

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