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MARBOCYL 100 MG/ML, SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Ima dovoljenje za promet

- Marbofloxacin

Informacije o zdravilu

Ime zdravila:
MARBOCYL 100 MG/ML, SOLUTION FOR INJECTION FOR CATTLE AND PIGS

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

Pot uporabe:
intramuskularna uporaba
Subkutana 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:

intramuskularna uporaba:

- **govedo**
  - Meat and offal. 6 day

Indication : Respiratory Dosage : 2 m/kg for 3 to 5 days (IV/IM/SC)

  - Milk. 36 hour

Indication : Mastitis Dosage : 2mg/kg for 3 days (IV/IM/SC)

  - Meat and offal. 6 day

Indication : Mastitis Dosage : 2mg/kg for 3 days (IV/IM/SC)

  - Milk. 72 hour

Indication : Respiratory Dosage : 8 mg/kg on a single occasion (IM)

  - Meat and offal. 3 day

Indication : Respiratory Dosage : 8 mg/kg on a single occasion (IM)

  - Milk. 36 hour

Indication : Respiratory Dosage : 2 m/kg for 3 to 5 days (IV/IM/SC)

- **Pig**
  - Meat and offal. 4 day

Subkutana uporaba:

- **govedo**
  - Meat and offal. 6 day

Indication : Respiratory Dosage : 2 m/kg for 3 to 5 days (IV/IM/SC)

- Milk. 36 hour

Indication : Mastitis Dosage : 2mg/kg for 3 days (IV/IM/SC)

- Meat and offal. 6 day

Indication : Mastitis Dosage : 2mg/kg for 3 days (IV/IM/SC)

- Milk. 72 hour

Indication : Respiratory Dosage : 8 mg/kg on a single occasion (IM)

- Meat and offal. 3 day

Indication : Respiratory Dosage : 8 mg/kg on a single occasion (IM)

- Milk. 36 hour

Indication : Respiratory Dosage : 2 m/kg for 3 to 5 days (IV/IM/SC)

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

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Na voljo v:

Portugal

Opis ovojnine zdravila:

Na voljo samo v English

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 Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Vetoquinol S.A.

Datum dovoljenja za promet z zdravilom:
26/10/1998

Proizvodna mesta, odgovorna za sproščanje serij:
Vetoquinol Biowet Sp. z o.o.
Vetoquinol S.A.

Pristojni organ:
Directorate General For Food And Veterinary

Številka dovoljenja za promet z zdravilom:
51247

Datum spremembe statusa dovoljenja za promet:
10/07/2025

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

Številka postopka:
FR/V/0107/001

Zadevne države članice:

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.