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Marbosol 100 mg/ml solution for injection for cattle and pigs

Ima dovoljenje za promet

- Marbofloxacin

Informacije o zdravilu

Ime zdravila:
Marbosol 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:
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 Multiple dose (2 mg/kg single daily injection, for 3 to 5 days)
  - Meat and offal. 6 day

Multiple dose (2 mg/kg single daily injection, for 3 to 5 days)

Subkutana uporaba:

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

Multiple dose (2 mg/kg single daily injection, for 3 to 5 days)

  - Milk. 36 hour Multiple dose (2 mg/kg single daily injection, for 3 to 5 days)

intramuskularna uporaba:

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

- **govedo**
  - Meat and offal. 3 day Single dose (8 mg/kg)
  - Meat and offal. 6 day

Multiple dose (2 mg/kg single daily injection, for 3 to 5 days)

  - Milk. 72 hour Single dose (8 mg/kg)

- Milk. 36 hour	Multiple dose (2 mg/kg single daily injection, for 3 to 5 days)

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

Na voljo v:
Germany

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:
CP-Pharma Handelsgesellschaft mbH

Datum dovoljenja za promet z zdravilom:
19/01/2013

Proizvodna mesta, odgovorna za sproščanje serij:
CP-Pharma Handelsgesellschaft mbH

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
401633.00.00

Datum spremembe statusa dovoljenja za promet:
10/08/2018

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

Številka postopka:
DE/V/0175/002

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 Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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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