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Fatroximin 2,94 mg/g uz ādas lietojams aerosols, šķīdums liellopiem, aitām, kazām, cūkām, zirgiem, trušiem, suņiem un kaķiem

Ima dovoljenje za promet

- Rifaximin

Informacije o zdravilu

Ime zdravila:
Fatroximin 2,94 mg/g uz ādas lietojams aerosols, šķīdums liellopiem, aitām, kazām, cūkām, zirgiem, trušiem, suņiem un kaķiem

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
Na voljo samo v Bulgarian Spanish Czech Danish Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
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
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Pot uporabe:
Dermalna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
2.94 milligram(s) / 1.00 gram(s)

Farmacevtska oblika:
Dermalno pršilo, raztopina

Karenca glede na pot uporabe:
Dermalna uporaba:

- **Rabbit**
  - Meat and offal. 0 day
- **govedo**

- Milk. 0 day
- Meat and offal. 0 day

- **Sheep**
  - Milk. 0 day
  - Meat and offal. 0 day

- **Goat**
  - Milk. 0 day
  - Meat and offal. 0 day

- **Horse**
  - Milk. 0 day
  - Meat and offal. 0 day

- **Pig**
  - Meat and offal. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QD06AX11

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:
Latvia

Opis ovojnine zdravila:
Na voljo samo v Latvian

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:
Fatro S.p.A.

Datum dovoljenja za promet z zdravilom:
5/07/2001

Proizvodna mesta, odgovorna za sproščanje serij:
Fatro S.p.A.

Pristojni organ:
Food And Veterinary Service

Številka dovoljenja za promet z zdravilom:
V/NRP/01/1370

Datum spremembe statusa dovoljenja za promet:
5/07/2001

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.