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Zactran 150 mg/ml - Solution for injection (cattle, pigs, sheep)

Ima dovoljenje za promet

- Gamithromycin

Informacije o zdravilu

Ime zdravila:
Zactran 150 mg/ml - Solution for injection (cattle, pigs, sheep)

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

Pot uporabe:
intramuskularna uporaba
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
150.00 milligram(s) / 1.00 Viala

Farmacevtska oblika:

Raztopina za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

- **govedo**
  - Meat and offal. 64 day — 64 days
- **Sheep**
  - Meat and offal. 29 day — 29 days
- **Pig**
  - Meat and offal. 16 day — 16 days

Subkutana uporaba:

- **govedo**
  - Meat and offal. 64 day — 64 days
- **Sheep**
  - Meat and offal. 29 day — 29 days
- **Pig**

- Meat and offal. 16 day	16 days

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

QJ01FA95

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:
Belgium , France , Germany , Luxembourg , Poland , Spain

Opis ovojnine zdravila:

Na voljo samo v English

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

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Pravna podlaga za izdajo dovoljenja za promet z zdravilom:

Na voljo samo v English Italian

Imetnik dovoljenja za promet z zdravilom:

Boehringer Ingelheim Vetmedica GmbH

Datum dovoljenja za promet z zdravilom:
24/07/2008

Proizvodna mesta, odgovorna za sproščanje serij:
Fresenius Kabi Austria GmbH
Boehringer Ingelheim Animal Health France

Pristojni organ:
European Commission

Številka dovoljenja za promet z zdravilom:
Ta podatek za to zdravilo ni na voljo.

Datum spremembe statusa dovoljenja za promet:
24/07/2008

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Combined File of all Documents

slovenščina (PDF)
Objavljeno na dan: 13/03/2026
Prenesi

ema-puar-zactran-v-129-var-x-0034-en.pdf

ema-puar-zactran-v-129-var-x-0027-en.pdf

ema-puar-zactran-v-129-par-en.pdf

ema-puar-zactran-v-129-var-ii-0036-en.pdf