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BACOLAM 100 mg/ml + 250.000 UI/ml sospensione iniettabile per bovini, ovi-caprini, suini, equini e tacchini

Ima dovoljenje za promet

- Amoxicillin
- COLISTIN SULFATE

Informacije o zdravilu

Ime zdravila:
BACOLAM 100 mg/ml + 250.000 UI/ml sospensione iniettabile per bovini, ovi-caprini, suini, equini e tacchini

Učinkovina:
Na voljo samo v English
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

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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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:
intramuskularna uporaba
Subkutana uporaba

Podatki o zdravilu

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

Na voljo samo v English
250000.00 international unit(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Suspenzija za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **govedo**
  - Meat and offal. 28 day

Non somministrare in animali in lattazione il cui latte o derivati siano destinati al consumo umano

- **Goat**

- Meat and offal. 28 day

Non somministrare in animali in lattazione il cui latte o derivati siano destinati al consumo umano

- **Sheep**
  - Meat and offal. 28 day

Non somministrare in animali in lattazione il cui latte o derivati siano destinati al consumo umano

- **Pig**
  - Meat and offal. 13 day

Subkutana uporaba:

- **Turkey**
  - Meat and offal. 25 day

Uso non consentito in animali che producono uova destinati al consumo umano

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Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QJ01RA01

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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Opis ovojnine zdravila:
Na voljo samo v Italian
Na voljo samo v Italian

Na voljo samo v Italian

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

Imetnik dovoljenja za promet z zdravilom:
Fatro S.p.A.

Datum dovoljenja za promet z zdravilom:
14/10/1993

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

Pristojni organ:
Ministry Of Health

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

Datum spremembe statusa dovoljenja za promet:
25/10/2013

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.