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IVERTIN 10 mg/ml Solution for Injection for Cattle and Pigs

Ima dovoljenje za promet

- Ivermectin

Informacije o zdravilu

Ime zdravila:
IVERTIN 10 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:
Subkutana uporaba

Podatki o zdravilu

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

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:
Subkutana uporaba:

- **govedo**
  - Milk. no withdrawal period

Do not use in lactating dairy cows producing milk for human consumption. Do not use in non-lactating dairy cows including pregnant heifers within 60 days of calving.

  - Meat and offal. 49 day

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

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QP54AA01

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 English
Na voljo samo v English
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Na voljo samo v English

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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 Norwegian

Imetnik dovoljenja za promet z zdravilom:
Laboratorios Calier S.A.

Datum dovoljenja za promet z zdravilom:
30/07/2013

Proizvodna mesta, odgovorna za sproščanje serij:
Laboratorios Calier S.A.

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
401914.00.00

Datum spremembe statusa dovoljenja za promet:
3/09/2018

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

Številka postopka:

DE/V/0165/001

Zadevne države članice:

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

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

Na voljo samo v Estonian English French Lithuanian Portuguese 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.