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Ecomectin 10 mg/ml Solution for Injection

Ima dovoljenje za promet

- Ivermectin

Informacije o zdravilu

Ime zdravila:
Ecomectin 10 mg/ml Solution for Injection

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:
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**
  - Meat and offal. 49 day
  - Milk. no withdrawal period

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

- **Sheep**
  - Meat and offal. 42 day
  - Milk. no withdrawal period

Do not use in lactating ewes producing milk for human consumption. Do not use in sheep which are intended to produce milk for human consumption within 60 days of lambing.

- **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

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
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:
Eco Animal Health Europe Limited

Datum dovoljenja za promet z zdravilom:
5/02/2004

Proizvodna mesta, odgovorna za sproščanje serij:
Produlab Pharma B.V.
Divasa Farmavic S.A.

Pristojni organ:
French Agency For Food, Environmental And Occupational Health & Safety

Številka dovoljenja za promet z zdravilom:
FR/V/4934184 5/2004

Datum spremembe statusa dovoljenja za promet:
5/02/2009

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

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Številka postopka:
IE/V/0144/001

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 Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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

Dokumenti

Package Leaflet and Labelling

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.