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Baycox Multi 50 mg/ml oral suspension for Cattle, Pigs and Sheep

Ima dovoljenje za promet

- Toltrazuril

Informacije o zdravilu

Ime zdravila:
Baycox Multi 50 mg/ml oral suspension for Cattle, Pigs and 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:
Peroralna uporaba

Podatki o zdravilu

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

Farmacevtska oblika:
Peroralna suspenzija

Karenca glede na pot uporabe:
Peroralna uporaba:

- **govedo**
  - Meat and offal. 63 day
  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

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

Not authorised for use in animals producing milk for human consumption.

- **Pig**
  - Meat and offal. 77 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QP51BC01

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

Na voljo v:
Norway

Opis ovojnine zdravila:
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:
Elanco Animal Health GmbH

Datum dovoljenja za promet z zdravilom:
29/06/2017

Proizvodna mesta, odgovorna za sproščanje serij:
KVP Pharma+Veterinaer Produkte GmbH

Pristojni organ:
Norwegian Medical Products Agency

Številka dovoljenja za promet z zdravilom:
15-10997

Datum spremembe statusa dovoljenja za promet:
21/09/2021

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/0360/001

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 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

Combined File of all Documents

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

Povzetek glavnih značilnosti zdravila

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

Navodilo za uporabo

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