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Enterisol Ileitis lyophilisate and diluent for oral suspension for pigs

Ima dovoljenje za promet

- Lawsonia intracellularis, strain MS B3903, Live

Informacije o zdravilu

Ime zdravila:
Enterisol Ileitis lyophilisate and diluent for oral suspension for pigs

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
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
Dajanje v krmno mešanico
Dajanje v vodo za pitje

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
79432.00 50% tissue culture infectious dose / 1.00 Dose

Farmacevtska oblika:

Liofilizat in vehikel za peroralno suspenzijo

Karenca glede na pot uporabe:

Peroralna uporaba:

- **Pig**
  - Meat and offal. 0 day

Dajanje v krmno mešanico:

- **Pig**
  - Meat and offal. 0 day

Dajanje v vodo za pitje:

- **Pig**
  - Meat and offal. 0 day

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

QI09AE04

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

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:
Boehringer Ingelheim Vetmedica GmbH

Datum dovoljenja za promet z zdravilom:
28/09/2008

Proizvodna mesta, odgovorna za sproščanje serij:
Boehringer Ingelheim Vetmedica GmbH

Pristojni organ:
Bulgarian Food Safety Authority

Številka dovoljenja za promet z zdravilom:
0022-1254

Datum spremembe statusa dovoljenja za promet:
28/09/2008

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

Številka postopka:
DE/V/0236/001

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

Povzetek glavnih značilnosti zdravila

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

Navodilo za uporabo

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Označevanje

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Combined File of all Documents

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