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Suiseng Coli /C Suspension for injection for pigs

Ima dovoljenje za promet

- Escherichia coli, LT toxoid
- Escherichia coli, fimbrial adhesin F5
- Escherichia coli, fimbrial adhesin F4ab
- Escherichia coli, fimbrial adhesin F6
- Escherichia coli, fimbrial adhesin F4ac
- Clostridium perfringens, type C, beta toxoid
- Clostridium novyi, type B, alpha toxoid

Informacije o zdravilu

Ime zdravila:
Suiseng Coli /C Suspension for injection for pigs

Učinkovina:
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
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:
intramuskularna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
55.00 percent / 1.00 Dose

Na voljo samo v English
79.00 percent / 1.00 Dose

Na voljo samo v English
65.00 percent / 1.00 Dose

Na voljo samo v English
80.00 percent / 1.00 Dose

Na voljo samo v English
78.00 percent / 1.00 Dose

Na voljo samo v English
1.05 relative potency / 1.00 Dose

Na voljo samo v English
1.23 relative potency / 1.00 Dose

Farmacevtska oblika:
Suspenzija za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

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

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI09AB08

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
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

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

Datum dovoljenja za promet z zdravilom:
1/04/2020

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

Pristojni organ:
Paul-Ehrlich-Institut

Številka dovoljenja za promet z zdravilom:
PEI.V.12026.01.1

Datum spremembe statusa dovoljenja za promet:
1/04/2020

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/0648/001

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

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.

Combined File of all Documents

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