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Porcilis ColiClos (--) - Suspension for injection

Ima dovoljenje za promet

- Escherichia coli, LT toxoid
- Clostridium perfringens, type C, strain 578, beta toxoid
- Escherichia coli, fimbrial adhesin F5
- Escherichia coli, fimbrial adhesin F6
- Escherichia coli, fimbrial adhesin F4ac
- Escherichia coli, fimbrial adhesin F4ab

Informacije o zdravilu

Ime zdravila:
Porcilis ColiClos (--) - Suspension for injection

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

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
Presentation_strength:≥10.9 log2 Ab titre Reference:Hse Index:0
Na voljo samo v English
Presentation_strength:≥20 IU Reference:Hse Index:1
Na voljo samo v English
Presentation_strength:≥8.4 log2 Ab titre Reference:Hse Index:2
Na voljo samo v English
Presentation_strength:≥7.8 log2 Ab titre Reference:Hse Index:3
Na voljo samo v English
Presentation_strength:≥8.1 log2 Ab titre Reference:Hse Index:4
Na voljo samo v English
Presentation_strength:≥9.7 log2 Ab titre Reference:Hse Index:5

Farmacevtska oblika:
Suspenzija za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **Pig**
  - Not applicable. 0 day Zero days

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

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

Na voljo v:
Austria , Belgium , Bulgaria , Croatia , Czechia , Denmark , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Lithuania , Netherlands , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Opis ovojnine zdravila:
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

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:
Intervet International B.V.

Datum dovoljenja za promet z zdravilom:
14/06/2012

Proizvodna mesta, odgovorna za sproščanje serij:
Intervet International B.V.

Pristojni organ:
European Commission

Številka dovoljenja za promet z zdravilom:
Ta podatek za to zdravilo ni na voljo.

Datum spremembe statusa dovoljenja za promet:
14/06/2012

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

Dokumenti

Combined File of all Documents

slovenščina (PDF)
Objavljeno na dan: 14/03/2026
Prenesi

ema-puar-v2011-porciliscoliclos-vra0018g-en.pdf

ema-puar-porcilic-coliclos-v-2011-par-en.pdf