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ENTERICOLIX, emulsion for injection for pigs

Ima dovoljenje za promet

- Escherichia coli, serotype O141:K85ab (fimbrial adhesin F6), strain P4, Inactivated
- Escherichia coli, serotype O101 (fimbrial adhesin F5 and F41), strain P10, Inactivated
- Escherichia coli, serotype O138:K81 (fimbrial adhesin F18ab), strain P5, Inactivated
- Escherichia coli, serotype O149:K91:H10 (fimbrial adhesin F4ac), strain P6, Inactivated
- Escherichia coli, serotype O157:H39 (fimbrial adhesin F18ac), strain P9, Inactivated
- Clostridium perfringens, type C, beta toxoid

Informacije o zdravilu

Ime zdravila:
ENTERICOLIX, emulsion 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

Ciljne živalske vrste:

Na voljo samo v Bulgarian Spanish Czech Danish German Estonian English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Danish German Estonian English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

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Pot uporabe:
intramuskularna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
1.00 relative potency / 2.00 millilitre(s)

Na voljo samo v English
1.00 relative potency / 2.00 millilitre(s)

Na voljo samo v English
1.00 relative potency / 2.00 millilitre(s)

Na voljo samo v English
1.00 relative potency / 2.00 millilitre(s)

Na voljo samo v English
1.00 relative potency / 2.00 millilitre(s)

Na voljo samo v English
10.00 Toxicity unit(s) / 2.00 millilitre(s)

Farmacevtska oblika:
Emulzija za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **Pig (sow, multipar)**
  - Meat and offal. 0 day

- **Pig (sow, primipar)**

- 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

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 Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
CZ Vaccines S.A.U.

Datum dovoljenja za promet z zdravilom:
8/03/2016

Proizvodna mesta, odgovorna za sproščanje serij:
CZ Vaccines S.A.U.

Pristojni organ:

Paul-Ehrlich-Institut

Številka dovoljenja za promet z zdravilom:
PEI.V.11739.01.1

Datum spremembe statusa dovoljenja za promet:
29/10/2020

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

Številka postopka:
ES/V/0228/001

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

Dokumenti

Označevanje

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.

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