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Salmoporc, lyophilisate and solvent for suspension for injection for pigs

Ima dovoljenje za promet

- Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 421/125 (histidine-adenine auxotrophic), Live

Informacije o zdravilu

Ime zdravila:
Salmoporc, lyophilisate and solvent for suspension for injection 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
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
8.00 log10 colony forming unit(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Liofilizat za suspenzijo za injiciranje

Karenca glede na pot uporabe:
Peroralna uporaba:

- **Pig**
  - Meat and offal. 6 week — six weeks after the second vaccination

Subkutana uporaba:

- **Pig**
  - Meat and offal. 6 week — six weeks after the second vaccination

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Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI09AE02

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 v:
Austria

Opis ovojnine zdravila:
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:
Ceva Sante Animale

Datum dovoljenja za promet z zdravilom:
25/04/2019

Proizvodna mesta, odgovorna za sproščanje serij:
Ceva-Phylaxia Zrt.
IDT Biologika GmbH

Pristojni organ:
Austrian Agency For Health And Food Safety

Številka dovoljenja za promet z zdravilom:
838869

Datum spremembe statusa dovoljenja za promet:
25/04/2019

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

Številka postopka:
NL/V/0247/001

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
Na voljo samo v Spanish Czech German Estonian English French Italian Latvian 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

Navodilo za uporabo

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.

Označevanje

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