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Covexin 10 Suspension for injection for sheep and cattle

Ima dovoljenje za promet

- Clostridium haemolyticum, toxoid
- Clostridium sordellii, toxoid
- Tetanus toxoid adsorbed
- Clostridium septicum, toxoid
- Clostridium novyi, toxoid
- Clostridium perfringens, type D, epsilon toxoid
- Clostridium perfringens, type B and C, beta toxoid
- Clostridium chauvoei, cells and toxin, Inactivated
- Clostridium perfringens, type A and C, alpha toxoid

Informacije o zdravilu

Ime zdravila:
Covexin 10 Suspension for injection for sheep and cattle

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
Na voljo samo v English
Na voljo samo v English

Ciljne živalske vrste:

govedo

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

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Pot uporabe:

Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
16.50 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Na voljo samo v English
0.80 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Na voljo samo v English
2.50 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Na voljo samo v English
3.60 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Na voljo samo v English
1.20 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Na voljo samo v English
5.10 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Na voljo samo v English
12.40 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Na voljo samo v English
90.00 percentage protection / 1.00 millilitre(s)

Na voljo samo v English
0.90 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Farmacevtska oblika:

Suspenzija za injiciranje

Karenca glede na pot uporabe:

Subkutana uporaba:

- **govedo**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI02AB01
QI04AB01

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:
Germany

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

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Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Zoetis Deutschland GmbH

Datum dovoljenja za promet z zdravilom:
24/09/2004

Proizvodna mesta, odgovorna za sproščanje serij:
Zoetis Belgium

Pristojni organ:
Paul-Ehrlich-Institut

Številka dovoljenja za promet z zdravilom:
PEI.V.03186.01.1

Datum spremembe statusa dovoljenja za promet:
28/10/2009

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

Številka postopka:
DE/V/0283/001

Zadevne države članice:
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

Combined File of all Documents

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