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Circovac (--) - Emulsion and suspension for emulsion for injection

Ima dovoljenje za promet

- Porcine circovirus type 2, strain 1010, Inactivated

Informacije o zdravilu

Ime zdravila:
Circovac (--) - Emulsion and suspension for emulsion for injection

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:
intramuskularna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
Presentation_strength:≥ 1.8 log 10 Elisa U Reference:Hse Index:0

Farmacevtska oblika:

Emulzija in suspenzija za emulzijo za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

- **Pig (female)**
  - Not applicable. 0 day Zero days

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

QI09AA07

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 , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Luxembourg , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

Opis ovojnine zdravila:
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:
Ceva-Phylaxia Zrt.

Datum dovoljenja za promet z zdravilom:
21/06/2007

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

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
21/06/2007

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-circovac-v-114-var-ii-0005-en.pdf

ema-puar-circovac-v-114-var-vra-0020-g-en.pdf

ema-puar-circovac-v-114-par-en.pdf

ema-puar-circovac-v-114-var-ws1945-en.pdf