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Coxevac (--) - Suspension for injection

Ima dovoljenje za promet

- Coxiella burnetii, strain Nine Mile, Inactivated

Informacije o zdravilu

Ime zdravila:
Coxevac (--) - Suspension 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
govedo
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:
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
Presentation_strength:≥72 QF unit Reference:Hse Index:0

Farmacevtska oblika:
Suspenzija za injiciranje

Karenca glede na pot uporabe:
Subkutana uporaba:

- **Sheep**
  - Meat and offal, milk. 0 day — Zero days
- **govedo**
  - Meat and offal, milk. 0 day — Zero days
- **Goat**
  - Meat and offal, milk. 0 day — Zero days

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI02AB

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

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Na voljo v:
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , France , Germany , Greece , Hungary , Ireland , Italy , Luxembourg , Netherlands , Poland , Portugal , Romania ,

Slovakia , Spain , United Kingdom (Northern Ireland)

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

Imetnik dovoljenja za promet z zdravilom:
Ceva Sante Animale

Datum dovoljenja za promet z zdravilom:
30/09/2010

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:
30/09/2010

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

Dokumenti

ema-puar-coxevac-v-155-par-en.pdf

ema-puar-coxevac-v-155-var-s-0007-en.pdf

ema-puar-coxevac-v-000155-var-x-0015-en.pdf