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Cevac Salmune ETI K (--) - Suspension for injection

Ima dovoljenje za promet

- Salmonella enterica, subsp. enterica, serovar Enteritidis, strain 038-90, Inactivated
- Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 076-94, Inactivated
- Salmonella enterica, subsp. enterica, serovar Infantis, strain SM-595, Inactivated

Informacije o zdravilu

Ime zdravila:
Cevac Salmune ETI K (--) - Suspension for injection

Učinkovina:
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 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:≥ 122 ELISA units Reference:In house Index:0
Na voljo samo v English
Presentation_strength:≥ 212 ELISA units Reference:In house Index:1
Na voljo samo v English
Presentation_strength:≥ 157 ELISA units Reference:In house Index:2

Farmacevtska oblika:
Suspenzija za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **Chicken**
  - Not applicable. 0 day Zero days

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI01AB01

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 samo v Spanish Czech German Estonian English French Italian 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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Na voljo v:
Belgium , Cyprus , Czechia , Germany , Greece , Hungary , 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 French Italian Latvian Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Ceva-Phylaxia Zrt.

Datum dovoljenja za promet z zdravilom:
30/08/2024

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:
30/08/2024

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-v6118-cevacsalmuneetik-en.pdf