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Cevac Meta L

Ima dovoljenje za promet

- Turkey rhinotracheitis virus, strain CRR126, Live

Informacije o zdravilu

Ime zdravila:
Cevac Meta L

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:
Okularna uporaba
Dajanje z razprševanjem

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
3.80 log10 50% tissue culture infectious dose / 1.00 Dose

Farmacevtska oblika:

Liofilizat za okulonazalno suspenzijo

Karenca glede na pot uporabe:

Okularna uporaba:

- **Chicken**
  - All relevant tissues. 0 day

Dajanje z razprševanjem:

- **Chicken**
  - All relevant tissues. 0 day

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

QI01AD01

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

Opis ovojnine zdravila:

Na voljo samo v English
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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 Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Ceva Sante Animale

Datum dovoljenja za promet z zdravilom:
22/07/2016

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

Pristojni organ:
French Agency For Food, Environmental And Occupational Health & Safety

Številka dovoljenja za promet z zdravilom:
FR/V/6805958 3/2016

Datum spremembe statusa dovoljenja za promet:
21/09/2021

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

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Številka postopka:
NL/V/0200/001

Zadevne države članice:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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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

Povzetek glavnih značilnosti zdravila

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