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CEVAC MASS L lyophilisate for oculonasal suspension for chickens

Ima dovoljenje za promet

- Infectious bronchitis virus, type Massachusetts, strain B-48, Live

Informacije o zdravilu

Ime zdravila:
CEVAC MASS L lyophilisate for oculonasal suspension for chickens

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
brojler

Pot uporabe:
Dajanje z razprševanjem

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
4.30 log 10 50% embryo infective dose / 1.00 Dose

Farmacevtska oblika:

Liofilizat za okulonazalno suspenzijo

Karenca glede na pot uporabe:

Dajanje z razprševanjem:

- **brojler**
  - Meat and offal. 0 day

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

QI01AD07

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

Na voljo v:

Denmark

Opis ovojnine zdravila:

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

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 Norwegian

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

Datum dovoljenja za promet z zdravilom:
28/06/2016

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

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
56876

Datum spremembe statusa dovoljenja za promet:
28/06/2016

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

Številka postopka:
HU/V/0125/001

Zadevne države članice:

Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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.

Combined File of all Documents

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