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CEVAC IBIRD LYOPHILISATE FOR OCULONASAL SUSPENSION/ USE IN DRINKING WATER FOR CHICKENS

Ima dovoljenje za promet

- Infectious bronchitis virus, type 793/B, strain 1/96, Live

Informacije o zdravilu

Ime zdravila:
CEVAC IBIRD LYOPHILISATE FOR OCULONASAL SUSPENSION/ USE IN DRINKING WATER FOR CHICKENS

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
brojler

Pot uporabe:
Okulonazalno dajanje
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
2.80 log 10 50% embryo infective dose / 1.00 Dose

Farmacevtska oblika:

Liofilizat za okulonazalno suspenzijo/dajanje v vodo za pitje

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Karenca glede na pot uporabe:

Okulonazalno dajanje:

- **Chicken (layer hen)**
  - All relevant tissues. 0 day

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

Peroralna uporaba:

- **Chicken (layer hen)**
  - All relevant tissues. 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

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Na voljo v:
Sweden

Opis ovojnine zdravila:
Na voljo samo v English
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:
13/05/2024

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

Pristojni organ:
Swedish Medical Products Agency

Številka dovoljenja za promet z zdravilom:
65060

Datum spremembe statusa dovoljenja za promet:
13/05/2024

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

Številka postopka:
FR/V/0245/001

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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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.

Navodilo za uporabo

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Označevanje

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

Combined File of all Documents

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