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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

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**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

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

QI01AD07

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**Pravni status za dobavo/izdajo zdravila:**

Zdravilo, ki se izdaja na veterinarski recept

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**Status dovoljenja za promet z zdravilom:**

Valid

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**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:**
Denmark

**Opis ovojnine zdravila:**
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
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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 Lithuanian Norwegian

**Imetnik dovoljenja za promet z zdravilom:**
Ceva Sante Animale

**Datum dovoljenja za promet z zdravilom:**
30/09/2013

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

**Pristojni organ:**
Danish Medicines Agency

**Številka dovoljenja za promet z zdravilom:**
51056

**Datum spremembe statusa dovoljenja za promet:**
30/09/2013

**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.

Combined File of all Documents

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

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