File downloaded on 2026-11-09
Source URL: https://medicines.health.europa.eu/veterinary/en/600000032282

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

Authorised

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

Product identification

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

Active substance:
Infectious bronchitis virus, type 793/B, strain 1/96, Live

Target species:
Chicken (layer hen)
Chicken (broiler)

Route of administration:
Oculonasal use
Oral use

Product details

Active substance and strength:
Infectious bronchitis virus, type 793/B, strain 1/96, Live
2.80 log 10 50% embryo infective dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate for oculonasal suspension/use in drinking water

Withdrawal period by route of administration:

Oculonasal use:

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

- **Chicken (broiler)**
  - All relevant tissues. 0 day

Oral use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD07

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Denmark

Available in:

Denmark

Package description:
Cardboard box of 1 bottle of 500 doses
Cardboard box of 20 bottles of 10000 doses
Cardboard box of 20 bottles of 5000 doses
Cardboard box of 20 bottles of 2500 doses
Cardboard box of 20 bottles of 1000 doses
Cardboard box of 20 bottles of 500 doses
Cardboard box of 10 bottles of 10000 doses
Cardboard box of 10 bottles of 5000 doses
Cardboard box of 10 bottles of 2500 doses
Cardboard box of 10 bottles of 1000 doses
Cardboard box of 10 bottles of 500 doses
Cardboard box of 1 bottle of 10000 doses
Cardboard box of 1 bottle of 5000 doses
Cardboard box of 1 bottle of 2500 doses
Cardboard box of 1 bottle of 1000 doses

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
30/09/2013

Manufacturing sites for batch release:
Ceva-Phylaxia Zrt.

Responsible authority:
Danish Medicines Agency

Authorisation number:
51056

Date of authorisation status change:
30/09/2013

Reference member state:
France

Procedure number:
FR/V/0245/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Malta Netherlands Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

eu-puar-frv0245001-mr-rpe774-en.pdf