CEVAC IBIRD LYOPHILISATE FOR SUSPENSION FOR CHICKENS

Authorised

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

Product identification

Medicine name:
CEVAC IBIRD LYOPHILISATE FOR SUSPENSION FOR CHICKENS
Cevac Ibird Lyophilisat pour suspension

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

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

Route of administration:
Oculonasal use
Oral use

Product details

Active substance and strength:
Avian 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 (broiler)**

  - All relevant tissues. 0 day

- **Chicken (layer hen)**

  - All relevant tissues. 0 day

- **Chicken (for reproduction)**

  - All relevant tissues. 0 day

Oral use:

- **Chicken (layer hen)**

  - All relevant tissues. 0 day

- **Chicken (for reproduction)**

  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD07

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Luxembourg

Package description:

Available only in French
Available only in French
Available only in French
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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:
29/05/2013

Manufacturing sites for batch release:
Ceva-Phylaxia Veterinary Biologicals Co. Ltd.

Responsible authority:
Le Ministere De La Sante Division De La Pharmacie Et Des Medicaments

Authorisation number:
V/915/17/04/1608

Date of authorisation status change:
10/02/2018

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

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