Avishield IB QX lyophilisate for oculonasal suspension/use in drinking water for chickens

Authorised

- Avian infectious bronchitis virus, type QX, strain 1285, Live

Product identification

Medicine name:
Avishield IB QX lyophilisate for oculonasal suspension/use in drinking water for chickens
Avishield IB QX liofilizat za okulonazalno suspenzijo/dajanje v vodo za pitje za piščance

Active substance:
Avian infectious bronchitis virus, type QX, strain 1285, Live

Target species:
Chicken

Route of administration:
Oculonasal use
In drinking water use

Product details

Active substance and strength:

Avian infectious bronchitis virus, type QX, strain 1285, Live
3.70 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**
  - Meat and offal. 0 day

In drinking water use:

- **Chicken**
  - Meat and offal. 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:
Slovenia

Package description:
Cardboard box with 10 vials of 1000 doses of vaccine
Cardboard box with 10 vials of 2500 doses of vaccine
Cardboard box with 10 vials of 5000 doses of vaccine

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Genera d.d.

Marketing authorisation date:
10/09/2025

Manufacturing sites for batch release:
Genera d.d.

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0826/001

Date of authorisation status change:
22/05/2025

Reference member state:
Netherlands

Procedure number:
NL/V/0429/001

Concerned member states:
Austria Belgium Croatia Czechia Denmark Estonia France Germany Greece Hungary Ireland Italy Latvia Lithuania Poland Portugal Romania Slovakia Slovenia Spain 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.

Labelling

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

Package Leaflet

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