

# Avishield IBD INT, lyophilisate for ocular nasal suspension/use in drinking water for chickens

Authorised

- Infectious bursal disease virus, strain VMG 91 (intermediate), Live

## Product identification

**Medicine name:**

Avishield IBD INT, lyophilisate for ocular nasal suspension/use in drinking water for chickens

---

**Active substance:**

Infectious bursal disease virus, strain VMG 91 (intermediate), Live

---

**Target species:**

Chicken

---

**Route of administration:**

Ocular nasal use  
Oral use

---

## Product details

**Active substance and strength:**

Infectious bursal disease virus, strain VMG 91 (intermediate), Live

4.00 log<sub>10</sub> 50% tissue culture infectious dose / 1.00 Dose

---

**Pharmaceutical form:**

Lyophilisate for ocular/nasal suspension/use in drinking water

---

**Withdrawal period by route of administration:**

**Ocular/nasal use:**

- 

**Chicken**

- All relevant tissues. 0 day

**Oral use:**

- 

**Chicken**

- All relevant tissues. 0 day

---

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01AD09

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Austria

---

**Package description:**

Carton box with 10 vials of 1000 doses of vaccine

Carton box with 10 vials of 2500 doses of vaccine

Carton box with 10 vials of 5000 doses of vaccine

---

## 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:**

Izo S.r.l.

---

**Marketing authorisation date:**

5/02/2019

---

**Manufacturing sites for batch release:**

Genera d.d.

---

**Responsible authority:**

Austrian Agency For Health And Food Safety

---

**Authorisation number:**

838682

---

**Date of authorisation status change:**

5/02/2019

---

**Reference member state:**

Netherlands

---

**Procedure number:**

NL/V/0244/001

---

**Concerned member states:**

Austria Belgium Czechia Germany Greece Hungary Ireland Italy Poland  
Portugal Romania Slovakia Spain

---

To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Package Leaflet

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

### 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.

### Combined File of all Documents