

# Avishield ND B1, Lyophilisate for Oculonasal Suspension/Use in Drinking Water for Chickens

Authorised

- Newcastle disease virus, strain B1 Hitchner, Live

## Product identification

**Medicine name:**

Avishield ND B1, Lyophilisate for Oculonasal Suspension/Use in Drinking Water for Chickens

---

**Active substance:**

Newcastle disease virus, strain B1 Hitchner, Live

---

**Target species:**

Chicken

---

**Route of administration:**

Oculonasal use  
Oral use

---

## Product details

**Active substance and strength:**

Newcastle disease virus, strain B1 Hitchner, Live  
6.00 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**

- Meat and offal. 0 day

**Oral use:**

- 

**Chicken**

- Meat and offal. 0 day

---

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

QI01AD06

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

United Kingdom (Northern Ireland)

---

**Package description:**

Carton with 10 vials of 2500 doses of vaccine. Colourless glass vials (type I), which are closed with brombutyl rubber stoppers and sealed with aluminium caps.

Carton with 10 vials of 5000 doses of vaccine. Colourless glass vials (type I), which are closed with brombutyl rubber stoppers and sealed with aluminium caps.

Carton with 10 vials of 1000 doses of vaccine. Colourless glass vials (type I), which are closed with brombutyl rubber stoppers and sealed with aluminium caps.

---

## 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:**

29/03/2018

---

**Manufacturing sites for batch release:**

Genera d.d.

---

**Responsible authority:**

The Veterinary Medicines Directorate

---

**Authorisation number:**

Vm 61812/3001

---

**Date of authorisation status change:**

18/04/2023

---

**Reference member state:**

Netherlands

---

**Procedure number:**

NL/V/0293/001

---

**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](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents