

# Avishield ND lyophilisate for oculonasal suspension/use in drinking water for chickens and turkeys

Authorised

- Newcastle disease virus, strain La Sota, Live

## Product identification

**Medicine name:**

Avishield ND lyophilisate for oculonasal suspension/use in drinking water for chickens and turkeys

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**Active substance:**

Newcastle disease virus, strain La Sota, Live

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**Target species:**

Turkey  
Chicken

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**Route of administration:**

Oculonasal use  
Oral use

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## Product details

### **Active substance and strength:**

Newcastle disease virus, strain La Sota, Live  
6.00 log<sub>10</sub> 50% tissue culture infectious dose / 1.00 Dose

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### **Pharmaceutical form:**

Lyophilisate for ocular nasal suspension/use in drinking water

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### **Withdrawal period by route of administration:**

#### **Ocular nasal use:**

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#### **Turkey**

- All relevant tissues. 0 day

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#### **Chicken**

- All relevant tissues. 1 day

#### **Oral use:**

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#### **Turkey**

- All relevant tissues. 0 day

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#### **Chicken**

- All relevant tissues. 1 day

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

QI01AD06

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### **Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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### **Authorisation status:**

Valid

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**Authorised in:**

Greece

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**Package description:**

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

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

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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Izo S.r.l.

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**Marketing authorisation date:**

19/09/2016

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**Manufacturing sites for batch release:**

Genera d.d.

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**Responsible authority:**

National Organization For Medicines

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**Authorisation number:**

74777/04-08-2021/K-0211201

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**Date of authorisation status change:**

3/08/2021

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0300/001

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**Concerned member states:**

Belgium Croatia Germany Greece Hungary Poland Portugal Romania  
Slovenia Spain United Kingdom (Northern Ireland)

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

### Package Leaflet

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

### Combined File of all Documents