

# AVINEW

Authorised

- Newcastle disease virus, strain VG/GA, Live

## Product identification

**Medicine name:**

AVINEW

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**Active substance:**

Newcastle disease virus, strain VG/GA, Live

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**Target species:**

Chicken (layer hen)

Chicken (for reproduction)

Chicken (broiler)

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**Route of administration:**

Ocular use

Oral use

Oculonasal use

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## Product details

**Active substance and strength:**

Newcastle disease virus, strain VG/GA, Live

5.50 log<sub>10</sub> 50% embryo infective dose / 1.00 Dose

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**Pharmaceutical form:**

Lyophilisate for suspension

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**Withdrawal period by route of administration:**

**Ocular use:**

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**Chicken (layer hen)**

- All relevant tissues. 0 day

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**Chicken (for reproduction)**

- All relevant tissues. 0 day

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**Chicken (broiler)**

- All relevant tissues. 0 day

**Oral use:**

•

**Chicken (layer hen)**

- All relevant tissues. 0 day

•

**Chicken (for reproduction)**

- All relevant tissues. 0 day

•

**Chicken (broiler)**

- All relevant tissues. 0 day

**Oculonasal use:**

•

**Chicken (layer hen)**

- All relevant tissues. 0 day

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**Chicken (for reproduction)**

- All relevant tissues. 0 day

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## **Chicken (broiler)**

- All relevant tissues. 0 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 except for some pack sizes

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### **Authorisation status:**

Valid

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### **Authorised in:**

France

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### **Package description:**

Box of one 2,000-dose bottle  
Box of ten 1,000-dose bottles  
Box of ten 2,000-dose bottles  
Box of one 1,000-dose bottle

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## **Additional information**

### **Entitlement type:**

Marketing Authorisation

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### **Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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### **Marketing authorisation holder:**

Boehringer Ingelheim Animal Health France

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### **Marketing authorisation date:**

13/12/1999

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### **Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France SCS

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/8349895 2/1999

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**Date of authorisation status change:**

13/12/2009

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**Reference member state:**

France

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**Procedure number:**

FR/V/0123/001

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**Concerned member states:**

Belgium Germany Greece Luxembourg Netherlands Portugal

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.