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# **AVINEW**

Not authorised

• Newcastle disease virus, strain VG/GA, Live

## **Product identification**

#### **Medicine name:**

**AVINEW** 

Avinew

#### **Active substance:**

Newcastle disease virus, strain VG/GA, Live

### **Target species:**

Chicken (layer hen)

Chicken (for reproduction)

Chicken (broiler)

#### Route of administration:

Ocular use

Oral use

Oculonasal use

### **Product details**

## **Active substance and strength:**

Newcastle disease virus, strain VG/GA, Live 5.50 log 10 50% embryo infective dose / 1.00 Dose

#### **Pharmaceutical form:**

Lyophilisate for suspension

### Withdrawal period by route of administration:

#### Ocular use:

•

### Chicken (layer hen)

- All relevant tissues. 0 day

•

### **Chicken (for reproduction)**

- All relevant tissues. 0 day

•

#### **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

•

### **Chicken (for reproduction)**

- All relevant tissues. 0 day

### Chicken (broiler)

- All relevant tissues. 0 day

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

**OI01AD06** 

### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Surrendered

#### **Authorised in:**

Germany

#### 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

## Additional information

### **Entitlement type:**

Marketing Authorisation

### Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

### Marketing authorisation holder:

Boehringer Ingelheim Vetmedica GmbH

## Marketing authorisation date:

5/03/2002

### Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority: Paul-Ehrlich-Institut
Authorisation number:
PEI.V.02053.01.1
Date of authorisation status change: 1/07/2025
Reference member state: France
Procedure number:
FR/V/0123/001
To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>
Documents

Summary of Product Characteristics

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