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# AVISAN CLON

Authorised

- Newcastle disease virus, strain CLON CL/79, Live

## Product identification

**Medicine name:**

AVISAN CLON

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**Active substance:**

Newcastle disease virus, strain CLON CL/79, Live

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**Target species:**

Chicken (broiler)

Chicken (pullet for egg production, future layer)

Future breeder pullet

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**Route of administration:**

Oculonasal use

In drinking water use

Nebulisation use

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## Product details

**Active substance and strength:**

Newcastle disease virus, strain CLON CL/79, Live

7.70 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:**

**Oculonasal use:**

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**Chicken (broiler)**

- Meat and offal. 0 day

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**Chicken (pullet for egg production, future layer)**

- Meat and offal. 0 day

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**Future breeder pullet**

- Meat and offal. 0 day

**In drinking water use:**

•

**Chicken (broiler)**

- Meat and offal. 0 day

•

**Chicken (pullet for egg production, future layer)**

- Meat and offal. 0 day

•

**Future breeder pullet**

- Meat and offal. 0 day

**Nebulisation use:**

•

**Chicken (broiler)**

- Meat and offal. 0 day

•

**Chicken (pullet for egg production, future layer)**

- Meat and offal. 0 day

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## **Future breeder pullet**

- Meat and offal. 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

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### **Authorisation status:**

Valid

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### **Authorised in:**

Spain

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### **Package description:**

Available only in [Spanish](#)

Available only in [Spanish](#)

Available only in [Spanish](#)

Available only in [Spanish](#)

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## Additional information

### **Entitlement type:**

Marketing Authorisation

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

Informed consent application (Article 13c of Directive No 2001/82/EC)

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### **Marketing authorisation holder:**

Laboratorios Hipra S.A.

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### **Marketing authorisation date:**

1/03/2021

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

Laboratorios Hipra S.A.

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**Responsible authority:**

Spanish Agency For Medicines And Medical Devices

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**Authorisation number:**

3974 ESP

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**Date of authorisation status change:**

2/03/2021

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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.

### Labelling

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