

# Hipraviar NDV-CLON Lyofilisaat voor suspensie voor oculonasaal gebruik/voor gebruik in drinkwater

Authorised

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

## Product identification

### **Medicine name:**

Hipraviar NDV-CLON Lyofilisaat voor suspensie voor oculonasaal gebruik/voor gebruik in drinkwater

### **Active substance:**

Newcastle disease virus, strain CLON CL/79, Live

### **Target species:**

Chicken (broiler)

### **Route of administration:**

Nebulisation use

In drinking water use

Oculonasal use

## Product details

### Active substance and strength:

Newcastle disease virus, strain CLON CL/79, Live  
6.50 log 10 50% embryo infective dose / 0.03 millilitre(s)

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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:

#### Nebulisation use:

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#### Chicken (broiler)

- Meat and offal. 0 day

#### In drinking water use:

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#### Chicken (broiler)

- Meat and offal. 0 day

#### Ocular nasal use:

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#### Chicken (broiler)

- 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:

Belgium

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**Package description:**

Hipraviar NDV-CLON 5 Vials with 1000 doses with Powder for solution and 5 Vials with 30 ml with solvent

Hipraviar NDV-CLON 10 Vials with 1000 doses with Powder for solution and 10 Vials with 30 ml with solvent

Hipraviar NDV-CLON 10 Vials with 1000 doses with Powder for solution and 10 Vials with 75 ml with solvent

Hipraviar NDV-CLON 10 Vials with 5000 doses with Powder for solution

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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:**

Laboratorios Hipra S.A.

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**Marketing authorisation date:**

24/06/1998

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**Manufacturing sites for batch release:**

Laboratorios Hipra S.A.

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**Responsible authority:**

Federal Agency For Medicines And Health Products

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**Authorisation number:**

BE-V193593

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**Date of authorisation status change:**

4/07/2024

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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.