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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)

Pharmaceutical form:
Lyophilisate for oculonasal suspension/use in drinking water

Withdrawal period by route of administration:

Nebulisation use:

- **Chicken (broiler)**
  - Meat and offal. 0 day

In drinking water use:

- **Chicken (broiler)**
  - Meat and offal. 0 day

Oculonasal use:

- **Chicken (broiler)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD06

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Laboratorios Hipra S.A.

Marketing authorisation date:
24/06/1998

Manufacturing sites for batch release:
Laboratorios Hipra S.A.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V193593

Date of authorisation status change:
4/07/2024

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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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.