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# Nobilis ND C2 lyophilisate for oculonasal suspension for chickens

Not authorised

- Newcastle disease virus, strain C2, Live

## Product identification

**Medicine name:**
Nobilis ND C2 lyophilisate for oculonasal suspension for chickens

**Active substance:**
Newcastle disease virus, strain C2, Live

**Target species:**
Chicken

**Route of administration:**
Oculonasal use
Nebulisation use

## Product details

**Active substance and strength:**
Newcastle disease virus, strain C2, Live
5.70 50% Embryo Infective Dose / 1.00 Dose

**Pharmaceutical form:**
Lyophilisate for oculonasal suspension

**Withdrawal period by route of administration:**

**Oculonasal use:**

- **Chicken**
  - All relevant tissues. no withdrawal period 0 days

**Nebulisation use:**

- **Chicken**
  - All relevant tissues. no withdrawal period 0 days

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QI01AD06

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

**Authorisation status:**
Surrendered

**Authorised in:**
Ireland

**Package description:**
Cardboard box with 1 vial of 500 doses.
Cardboard box with 1 vial of 5000 doses.
PET plastic boxes with 12 cups of 5000 doses
PET plastic boxes with 12 cups of 2500 doses
PET plastic boxes with 12 cups of 10000 doses
PET plastic boxes with 12 cups of 1000 doses
Cardboard box with 1 vial of 25000 doses.
Cardboard box with 1 vial of 2500 doses.
Cardboard box with 1 vial of 10000 doses.
Cardboard box with 1 vial of 1000 doses.

Cardboard box with 10 vials of 1000 doses.
Cardboard box with 10 vials of 10000 doses.
Cardboard box with 10 vials of 2500 doses.
Cardboard box with 10 vials of 25000 doses.
Cardboard box with 10 vials of 5000 doses.
Cardboard box with 10 vials of 500 doses.

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Intervet (Ireland) Limited

**Marketing authorisation date:**
3/06/2005

**Manufacturing sites for batch release:**
Intervet International B.V.

**Responsible authority:**
Health Products Regulatory Authority

**Authorisation number:**
VPA10996/187/001

**Date of authorisation status change:**
31/10/2024

**Reference member state:**
Netherlands

**Procedure number:**
NL/V/0113/001

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

## Documents

Combined File of all Documents