

# Nobilis ND Clone 30, liofilizatas suspensijai į akį ar šnervę ruošti / naudoti su geriamuoju vandeniu vištoms ir kalakutams

Authorised

- Newcastle disease virus, strain Clone 30, Live

## Product identification

### Medicine name:

Nobilis ND Clone 30, liofilizatas suspensijai į akį ar šnervę ruošti / naudoti su geriamuoju vandeniu vištoms ir kalakutams

### Active substance:

Newcastle disease virus, strain Clone 30, Live

### Target species:

Chicken

Turkey

### Route of administration:

Oculonasal use

Nebulisation use

In drinking water use

## Product details

### Active substance and strength:

Newcastle disease virus, strain Clone 30, Live  
6.00 log 10 50% embryo lethal dose / 1.00 Dose

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### Pharmaceutical form:

Lyophilisate for oculonasal suspension/use in drinking water

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### Withdrawal period by route of administration:

#### Oculonasal use:

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##### Chicken

- Meat and offal. 0 day

- 

##### Turkey

- Meat and offal. 0 day

#### Nebulisation use:

- 

##### Chicken

- Meat and offal. 0 day

- 

##### Turkey

- Meat and offal. 0 day

#### In drinking water use:

- 

##### Chicken

- Meat and offal. 0 day

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##### Turkey

- Meat and offal. 0 day

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

QI01AD06

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**Legal status of supply:**

This information is not available for this product.

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**Authorisation status:**

Valid

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**Authorised in:**

Lithuania

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**Package description:**

Available only in [Lithuanian](#)

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Available only in [Lithuanian](#)

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Available only in [Lithuanian](#)

Available only in [Lithuanian](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis reviewed according to Acquis communautaire

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

21/12/1995

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

State Food And Veterinary Service

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**Authorisation number:**

LT/2/95/0259/001-007

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**Date of authorisation status change:**

9/01/2007

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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.

RV0259.pdf