

# Nobilis RT+IBmulti+ND+EDS emulsion for injection for chickens

Authorised

- Newcastle disease virus, strain Clone 30, Inactivated
- Eggdrop syndrome-1976 virus, strain BC14, Inactivated
- Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated
- Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

## Product identification

**Medicine name:**

Nobilis RT+IBmulti+ND+EDS emulsion for injection for chickens

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**Active substance:**

Newcastle disease virus, strain Clone 30, Inactivated

Eggdrop syndrome-1976 virus, strain BC14, Inactivated

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated

Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

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**Target species:**

Chicken

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**Route of administration:**

Intramuscular use

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## Product details

### **Active substance and strength:**

Newcastle disease virus, strain Clone 30, Inactivated  
50.00 50% Protective Dose / 0.50 millilitre(s)

Eggdrop syndrome-1976 virus, strain BC14, Inactivated  
6.50 log<sub>2</sub> haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated  
9.50 log<sub>2</sub> enzyme-linked immunosorbent assay unit(s) / 0.50 millilitre(s)

Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated  
4.00 log<sub>2</sub> virus neutralising unit(s) / 0.50 millilitre(s)

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated  
5.50 log<sub>2</sub> virus neutralising unit(s) / 0.50 millilitre(s)

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### **Pharmaceutical form:**

Emulsion for injection

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### **Withdrawal period by route of administration:**

#### **Intramuscular use:**

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#### **Chicken**

- Meat and offal. 0 day
  - Egg. 0 day
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### **Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01AA18

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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:**

Czechia

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**Available in:**

Czechia

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**Package description:**

(ID2) 500 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 500 millilitre(s)

(ID1) 250 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 250 millilitre(s)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

30/07/2004

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

97/127/04-C

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**Date of authorisation status change:**

17/09/2009

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0209/001

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**Concerned member states:**

Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia France Greece  
Ireland Italy Latvia Lithuania Luxembourg Netherlands Portugal Spain  
United Kingdom (Northern Ireland)

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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.

Combined File of all Documents