

# Nobilis RT+IBmulti+G+ND Emulsion for injection (water-in-oil)

Not  
authorised

- Newcastle disease virus, strain Clone 30, Inactivated
- Infectious bursal disease virus, strain D78, 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+G+ND Emulsion for injection (water-in-oil)

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**Active substance:**

Newcastle disease virus, strain Clone 30, Inactivated

Infectious bursal disease virus, strain D78, 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)

Infectious bursal disease virus, strain D78, Inactivated

14.50 log<sub>2</sub> virus neutralising 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:**

QI01AA06

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Surrendered

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**Authorised in:**

Italy

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**Package description:**

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

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

24/04/2000

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

103074

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**Date of authorisation status change:**

6/11/2025

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**Reference member state:**

Germany

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**Procedure number:**DE/V/0212/001

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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

Combined File of all Documents

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