

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

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)

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

Target species:

Chicken

Route of administration:

Intramuscular use

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 log₂ virus neutralising unit(s) / 0.50 millilitre(s)

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated

9.50 log₂ enzyme-linked immunosorbent assay unit(s) / 0.50 millilitre(s)

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

4.00 log₂ virus neutralising unit(s) / 0.50 millilitre(s)

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

5.50 log₂ virus neutralising unit(s) / 0.50 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:**Intramuscular use:**

-

Chicken

- Meat and offal. 0 day

- Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

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)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Intervet

Marketing authorisation date:

29/06/2000

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/9652212 5/2000

Date of authorisation status change:

29/06/2010

Reference member state:

Germany

Procedure number:

DE/V/0212/001

Concerned member states:

Austria Belgium Denmark France Greece Ireland Italy Luxembourg
Netherlands Portugal Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 28/01/2022

Updated on: 13/03/2026

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Package Leaflet and Labelling

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