Nobilis RT + IBmulti + G + ND

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

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

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Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Available in:
United Kingdom (Northern Ireland)

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)

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 International B.V.

Marketing authorisation date:
27/04/2000

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 06376/3012

Date of authorisation status change:
20/06/2024

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