

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

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

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)

Eggdrop syndrome-1976 virus, strain BC14, Inactivated

6.50 log₂ haemagglutination inhibiting 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:

QI01AA18

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Estonia

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:

4/05/2015

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

State Agency Of Medicines

Authorisation number:

1904

Date of authorisation status change:

4/05/2015

Reference member state:

Germany

Procedure number:

DE/V/0209/001

Concerned member states:

Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia France Greece
Ireland Italy Latvia Lithuania 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)

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