

NOBILIS RT+IBmulti+G+ND, injekcinė emulsija vištoms

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, injekcinė emulsija vištoms

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 (for reproduction)

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 (for reproduction)

- Meat and offal. 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:

Lithuania

Available in:

Lithuania

Package description:

Available only in [Lithuanian](#)

Available only in [Lithuanian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

23/04/2003

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/03/1555/001-002

Date of authorisation status change:

5/06/2023

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

Documents

Combined File of all Documents

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