Nobilis RT+IBmulti+G+ND emulsija injekcijām vistām

Authorised

- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
- Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated
- Infectious bursal disease virus, strain D78, Inactivated
- Newcastle disease virus, strain Clone 30, Inactivated

Product identification

Medicine name:
Nobilis RT+IBmulti+G+ND emulsija injekcijām vistām

Active substance:
Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated
Infectious bursal disease virus, strain D78, Inactivated
Newcastle disease virus, strain Clone 30, Inactivated

Target species:
Chicken (for reproduction)

Route of administration:

Intramuscular use

Product details

Active substance and strength:
Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
5.50 log2 virus neutralising unit(s) / 1.00 unit(s)

Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
4.00 log2 virus neutralising unit(s) / 1.00 unit(s)

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated
9.50 log2 enzyme-linked immunosorbent assay unit(s) / 1.00 unit(s)

Infectious bursal disease virus, strain D78, Inactivated
14.50 log2 virus neutralising unit(s) / 1.00 unit(s)

Newcastle disease virus, strain Clone 30, Inactivated
50.00 50% Protective Dose / 1.00 unit(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:
Latvia

Available in:
Latvia

Package description:
Available only in Latvian
Available only in Latvian

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:
14/02/2003

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

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/03/1535

Date of authorisation status change:
16/02/2003

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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