NOBILIS RT + IBmulti + ND + EDS

Authorised

- Eggdrop syndrome-1976 virus, strain V127, Inactivated
- Newcastle disease virus, strain Clone 30, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
- Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated

Product identification

Medicine name:
NOBILIS RT + IBmulti + ND + EDS

Active substance:
Eggdrop syndrome-1976 virus, strain V127, Inactivated

Newcastle disease virus, strain Clone 30, Inactivated

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

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

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated

Target species:
Chicken (for reproduction)

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:

Eggdrop syndrome-1976 virus, strain V127, Inactivated
6.50 log2 haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Newcastle disease virus, strain Clone 30, Inactivated
50.00 50% Protective Dose / 0.50 millilitre(s)

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
5.50 log2 virus neutralising 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)

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated
9.50 log2 enzyme-linked immunosorbent assay unit(s) / 0.50 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA18

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Romania

Available in:

Romania

Package description:

Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
20/04/2006

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

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
120265

Date of authorisation status change:
21/03/2024

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

Documents

Summary of Product Characteristics

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