

Nobilis RT+IB multi+G+ND vakcina A.U.V.

Authorised

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

Product identification

Medicine name:

Nobilis RT+IB multi+G+ND vakcina A.U.V.

Active substance:

Infectious bursal disease virus, strain D78, Inactivated

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated

Newcastle disease virus, strain Clone 30, Inactivated

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Target species:

Chicken (hen)

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Infectious bursal disease virus, strain D78, Inactivated

14.50 log₂ virus neutralising unit(s) / 1.00 unit(s)/dose

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated

9.50 log₂ enzyme-linked immunosorbent assay unit(s) / 1.00 unit(s)/dose

Newcastle disease virus, strain Clone 30, Inactivated

4.00 log₂ haemagglutination inhibiting unit(s) / 1.00 unit(s)/dose

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

4.00 log₂ virus neutralising unit(s) / 1.00 unit(s)/dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Chicken (hen)

- Meat and offal. 4 week

Subcutaneous use:

-

Chicken (hen)

- Meat and offal. 4 week

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Hungary

Available in:

Hungary

Package description:

Available only in Hungarian

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:

4/12/2001

Manufacturing sites for batch release:

INTERVET INTERNATIONAL B.V.

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

4/12/2001

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