

Nobilis REO+IB+G+ND vakcina A.U.V.

Authorised

- Avian reovirus, strain 2408, Inactivated
- Avian reovirus, strain 1733, Inactivated
- Infectious bursal disease virus, Inactivated
- Newcastle disease virus, Inactivated
- Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Product identification

Medicine name:

Nobilis REO+IB+G+ND vakcina A.U.V.

Active substance:

Avian reovirus, strain 2408, Inactivated

Avian reovirus, strain 1733, Inactivated

Infectious bursal disease virus, Inactivated

Newcastle disease virus, Inactivated

Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Target species:

Chicken (hen)

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Avian reovirus, strain 2408, Inactivated

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

Avian reovirus, strain 1733, Inactivated

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

Infectious bursal disease virus, Inactivated

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

Newcastle disease virus, Inactivated

50.00 50% Protective Dose / 1.00 unit(s)/dose

Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

6.00 log₂ haemagglutination inhibiting 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. 0 day

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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA16

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised 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:

13/05/1996

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:

13/05/1996

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