Nobilis IB+ND+EDS vakcina A.U.V.

Authorised

- Newcastle disease virus, strain Clone 30, Inactivated
- Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Egg drop syndrome '76 virus, strain BC14, Inactivated

Product identification

Medicine name:
Nobilis IB+ND+EDS vakcina A.U.V.

Active substance:
Newcastle disease virus, strain Clone 30, Inactivated

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

Egg drop syndrome '76 virus, strain BC14, Inactivated

Target species:
Chicken (hen)

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Newcastle disease virus, strain Clone 30, Inactivated

4.00 log2 haemagglutination inhibiting unit(s) / 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

Egg drop syndrome '76 virus, strain BC14, Inactivated
6.50 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
  - Meat and offal. 0 day

Subcutaneous use:

- **Chicken (hen)**
  - Meat and offal. 0 day
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AA13

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:
This information is not available for this product.

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:
8/09/1999

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

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