

Nobilis IB+ND+EDS

Authorised

- Newcastle disease virus, strain Clone 30, Inactivated
- Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Eggdrop syndrome-1976 virus, strain BC14, Inactivated

Product identification

Medicine name:

Nobilis IB+ND+EDS

Active substance:

Newcastle disease virus, strain Clone 30, Inactivated

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

Eggdrop syndrome-1976 virus, strain BC14, Inactivated

Target species:

Chicken

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Newcastle disease virus, strain Clone 30, Inactivated

50.00 50% Protective Dose / 0.50 millilitre(s)

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

6.00 log₂ haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Eggdrop syndrome-1976 virus, strain BC14, Inactivated

6.50 log₂ haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Chicken

- Egg. 0 day

Subcutaneous use:

-

Chicken

- Egg. 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:

Germany

Available in:

Germany

Package description:

Available only in German

Available only in German

Available only in [German](#)

Available only in [German](#)

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 Deutschland GmbH

Marketing authorisation date:

4/11/1996

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Paul-Ehrlich-Institut

Authorisation number:

492a/93

Date of authorisation status change:

1/06/2011

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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