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Nobilis IBmulti+ND+EDS Emulsion for injection (water-in-oil)

Authorised

- Newcastle disease virus, strain Clone 30, Inactivated
- Eggdrop syndrome-1976 virus, strain BC14, Inactivated
- Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Product identification

Medicine name:
Nobilis IBmulti+ND+EDS Emulsion for injection (water-in-oil)

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

Eggdrop syndrome-1976 virus, strain BC14, Inactivated

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

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Target species:
Chicken (for reproduction)

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

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

Eggdrop syndrome-1976 virus, strain BC14, Inactivated
6.50 log2 haemagglutination inhibiting 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)

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
5.50 log2 virus neutralising unit(s) / 0.50 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Chicken (for reproduction)**
  - Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA13

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:

(ID3): 1 Box with 1 Bottle (Glass) with 250 millilitre(s) (500 ID)
(ID4): 1 Box with 1 Bottle (Glass) with 500 millilitre(s) (1000 ID)
(ID2): 1 Box with 1 Bottle (PolyEthylene TerePhthalate) with 500 millilitre(s) (1000 ID)
(ID1): 1 Box with 1 Bottle (PolyEthylene TerePhthalate) with 250 millilitre(s) (500 ID)

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:
30/01/2006

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

Responsible authority:
Danish Medicines Agency

Authorisation number:
38233

Date of authorisation status change:
30/01/2006

Reference member state:
Germany

Procedure number:
DE/V/0225/001

Concerned member states:

Belgium Denmark France Ireland Luxembourg Sweden

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents