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

Authorised

- Newcastle disease virus, strain Clone 30, Inactivated
- Infectious bursal disease virus, strain D78, Inactivated
- Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated
- Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Product identification

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

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

Infectious bursal disease virus, strain D78, Inactivated

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated

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

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Target species:
Chicken

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Newcastle disease virus, strain Clone 30, Inactivated
50.00 50% Protective Dose / 0.50 millilitre(s)

Infectious bursal disease virus, strain D78, Inactivated
14.50 log2 virus neutralising unit(s) / 0.50 millilitre(s)

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated
9.50 log2 enzyme-linked immunosorbent assay 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:
Intramuscular use:

- **Chicken**
  - Meat and offal. 0 day
  - Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AA06

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Netherlands

Available in:
Netherlands

Package description:
(ID2) 500 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 500 millilitre(s)
(ID1) 250 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 250 millilitre(s)

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 Nederland B.V.

Marketing authorisation date:
8/07/2009

Manufacturing sites for batch release:
INTERVET INTERNATIONAL B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 9707

Date of authorisation status change:
8/02/2022

Reference member state:
Germany

Procedure number:
DE/V/0212/001

Concerned member states:
Austria Belgium Denmark France Greece Ireland Italy Luxembourg Netherlands Portugal Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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

Summary of Product Characteristics