Nobilis RT+IBmulti+ND+EDS emulsion for injection for chickens

Authorised

- Newcastle disease virus, strain Clone 30, Inactivated
- Eggdrop syndrome-1976 virus, strain BC14, 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+ND+EDS emulsion for injection for chickens

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

Eggdrop syndrome-1976 virus, strain BC14, 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)

Eggdrop syndrome-1976 virus, strain BC14, Inactivated
6.50 log2 haemagglutination inhibiting 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:
QI01AA18

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

Authorisation status:
Valid

Authorised in:

Ireland

Available in:
Ireland

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 (Ireland) Limited

Marketing authorisation date:
3/11/2004

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

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10996/181/001

Date of authorisation status change:
3/11/2004

Reference member state:
Germany

Procedure number:
DE/V/0209/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia France Greece Ireland Italy Latvia Lithuania 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