Nobilis RT + IBmulti + G + ND, Injekční emulze

Authorised

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

Product identification

Medicine name:
Nobilis RT + IBmulti + G + ND, Injekční emulze

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

Target species:
Chicken

Route of administration:

Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
5.50 log2 virus neutralising unit(s) / 1.00 Dose

Newcastle disease virus, strain Clone 30, Inactivated
4.00 log2 haemagglutination inhibiting unit(s) / 1.00 Dose

Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
4.00 log2 virus neutralising unit(s) / 1.00 Dose

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated
9.50 log2 enzyme-linked immunosorbent assay unit(s) / 1.00 Dose

Infectious bursal disease virus, strain D78, Inactivated
14.50 log2 virus neutralising unit(s) / 1.00 Dose

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Subcutaneous use:

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

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:
Czechia

Available in:
Czechia

Package description:
Available only in Czech
Available only in Czech

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
24/01/2000

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

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/015/00-C

Date of authorisation status change:
25/04/2014

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.

Package Leaflet

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

Labelling

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