Nobilis RT + IB multi + G + ND emulsja do wstrzykiwań dla kur

Authorised

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

Product identification

Medicine name:
Nobilis RT + IB multi + G + ND emulsja do wstrzykiwań dla kur

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

Target species:
Chicken (hen)

Route of administration:

Intramuscular use

Product details

Active substance and strength:
Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
1.00 unknown / 1.00 unknown

Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
1.00 unknown / 1.00 unknown

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated
1.00 unknown / 1.00 unknown

Infectious bursal disease virus, strain D78, Inactivated
1.00 unknown / 1.00 unknown

Newcastle disease virus, strain Clone 30, Inactivated
50.00 50% Protective Dose / 1.00 50% Protective Dose

Pharmaceutical form:
Emulsion for injection/infusion

Withdrawal period by route of administration:
Intramuscular use:

- **Chicken (hen)**
  - Meat and offal. 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:
Poland

Available in:
Poland

Package description:
Available only in Polish
Available only in Polish

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:
9/02/2001

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

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
1131

Date of authorisation status change:
9/02/2001

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

Documents

Labelling

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.

Summary of Product Characteristics

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