Nobilis RT + IB Multi + ND + EDS emulsja do wstrzykiwań dla kur

Authorised

- Newcastle disease virus, strain Clone 30, Inactivated
- Eggdrop syndrome-1976 virus, strain BC14, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
- Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated

Product identification

Medicine name:
Nobilis RT + IB Multi + ND + EDS emulsja do wstrzykiwań dla kur

Active substance:
Newcastle disease virus, strain Clone 30, Inactivated
Eggdrop syndrome-1976 virus, strain BC14, Inactivated
Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated

Target species:
Chicken (hen)

Route of administration:

Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Newcastle disease virus, strain Clone 30, Inactivated
50.00 50% Protective Dose / 1.00 50% Protective Dose

Eggdrop syndrome-1976 virus, strain BC14, Inactivated
1.00 unknown / 1.00 unknown

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

Pharmaceutical form:
Emulsion for injection/infusion

Withdrawal period by route of administration:
Subcutaneous use:

- **Chicken (hen)**
  - Meat and offal. 0 day

Intramuscular use:

- **Chicken (hen)**
  - Meat and offal. 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:
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.
MERCK SHARP & DOHME ANIMAL HEALTH, S.L.

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

Authorisation number:
1118

Date of authorisation status change:

9/02/2001

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

Documents

Package Leaflet

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Labelling

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.