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Nobilis RT + IB Multi + ND + EDS emulsja do wstrzykiwa? dla kur

- 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

Authorised

Product identification

Medicine name:
Nobilis RT + IB Multi + ND + EDS emulsja do wstrzykiwa? 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

Product information

Package Leaflet

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