Nobilis RT+Ibmulti+G+ND, emulzija za injekciju, za kokoši

Authorised

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

Product identification

Medicine name:
Nobilis RT+Ibmulti+G+ND, emulzija za injekciju, za kokoši

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

Target species:
Chicken (pullet for egg production, future layer)

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)

Infectious bursal disease virus, strain D78, Inactivated
14.50 log2 virus neutralising 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)

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated
9.50 log2 enzyme-linked immunosorbent assay unit(s) / 0.50 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Chicken (pullet for egg production, future layer)**
  - 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:
Croatia

Package description:
Available only in Croatian
Available only in Croatian

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. Subsidiary In The Republic Of Croatia

Marketing authorisation date:
23/07/2013

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

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/18-01/333

Date of authorisation status change:
3/06/2024

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

Documents

Summary of Product Characteristics

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