Tabic V.H. putojošā tablete vistām un tītariem

Authorised

- Newcastle disease virus, strain V.H., Live

Product identification

Medicine name:
Tabic V.H. putojošā tablete vistām un tītariem

Active substance:
Newcastle disease virus, strain V.H., Live

Target species:
Turkey
Chicken (broiler)
Chicken (for reproduction)
Chicken (layer hen)

Route of administration:
In drinking water use
Intraocular use
Nebulisation use

Product details

Active substance and strength:
Newcastle disease virus, strain V.H., Live

1000000.00 50% Embryo Infective Dose / 1.00 unit(s)

Pharmaceutical form:

Effervescent tablet

Withdrawal period by route of administration:

In drinking water use:

- **Turkey**
  - Not specified. 0 day
- **Chicken (broiler)**
  - Not specified. 0 day
- **Chicken (for reproduction)**
  - Not specified. 0 day
- **Chicken (layer hen)**
  - Not specified. 0 day

Intraocular use:

- **Chicken (for reproduction)**
  - Not specified. 0 day
- **Turkey**
  - Not specified. 0 day
- **Chicken (broiler)**
  - Not specified. 0 day
- **Chicken (layer hen)**
  - Not specified. 0 day

Nebulisation use:

- **Chicken (for reproduction)**
  - Not specified. 0 day
- **Chicken (layer hen)**
  - Not specified. 0 day
- **Turkey**
  - Not specified. 0 day
- **Chicken (broiler)**
  - Not specified. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD06

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:
Valid

Authorised in:
Latvia

Available in:
Latvia

Package description:
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian

Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Phibro Animal Health (Poland) Sp. z o.o.

Marketing authorisation date:
16/12/2010

Manufacturing sites for batch release:
Synoptis Industrial Sp. z o.o.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/10/0032

Date of authorisation status change:
16/12/2010

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

Documents

Combined File of all Documents

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