File downloaded on 2026-11-08
Source URL: https://medicines.health.europa.eu/veterinary/en/600000096501

TAbic V.H., šnypščioji tabletė vištoms ir kalakutams

Authorised

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

Product identification

Medicine name:
TAbic V.H., šnypščioji tabletė vištoms ir kalakutams

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

Target species:
Chicken
Turkey

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

Product details

Active substance and strength:
Newcastle disease virus, strain V.H., Live
1000000.00 50% Embryo Infective Dose / 1.00 Dose

Pharmaceutical form:
Effervescent tablet

Withdrawal period by route of administration:
In drinking water use:

- **Chicken**
  - Meat and offal. 0 day

- **Turkey**
  - Meat and offal. 0 day

Ocular use:

- **Chicken**
  - Meat and offal. 0 day

- **Turkey**
  - Meat and offal. 0 day

Nebulisation use:

- **Chicken**
  - Meat and offal. 0 day

- **Turkey**
  - Meat and offal. 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:
Lithuania

Available in:
Lithuania

Package description:
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian

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:
28/12/2008

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

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/08/1827/001-014

Date of authorisation status change:
17/12/2013

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.