V.H. + H-120 liofilizāts ūdens šķīduma pagatavošanai vistām

Authorised

- Newcastle disease virus, strain V.H., Live
- Infectious bronchitis virus, type Massachusetts, strain H120, Live

Product identification

Medicine name:
V.H. + H-120 liofilizāts ūdens šķīduma pagatavošanai vistām

Active substance:
Newcastle disease virus, strain V.H., Live
Infectious bronchitis virus, type Massachusetts, strain H120, Live

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

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

Product details

Active substance and strength:

Newcastle disease virus, strain V.H., Live

Infectious bronchitis virus, type Massachusetts, strain H120, Live

---

Pharmaceutical form:

Lyophilisate for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

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

Intraocular use:

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

Nebulisation use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD

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

Authorisation status:
Valid

Authorised in:
Latvia

Package description:
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/10/0033

Date of authorisation status change:
16/12/2010

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

Documents

Package Leaflet

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.

Labelling

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