# V.H. + H-120 - atenuowany wirus rzekomego pomoru ptaków (choroby Newcastle), szczep VHnie mniej niż 10^6,0 EID50 i nie więcej niż 10^7,49 EID50;- atenuowany wirus zakaźnego zapalenia oskrzeli ptaków, szczep H-120nie mniej niż 10^3,5 EID50 i nie więcej niż 10^4,49 EID50. Liofilizat do sporządzania zawiesiny

Authorised

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

## Product identification

**Medicine name:**
V.H. + H-120 - atenuowany wirus rzekomego pomoru ptaków (choroby Newcastle), szczep VHnie mniej niż 10^6,0 EID50 i nie więcej niż 10^7,49 EID50;- atenuowany

wirus zakaźnego zapalenia oskrzeli ptaków, szczep H-120nie mniej niż 10^3,5 EID50 i nie więcej niż 10^4,49 EID50. Liofilizat do sporządzania zawiesiny

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

**Target species:**
Chicken (hen)

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

## Product details

**Active substance and strength:**
Newcastle disease virus, strain V.H., Live
30903000.00 50% Embryo Infective Dose / 1.00 50% Embryo Infective Dose
Avian infectious bronchitis virus, type Massachusetts, strain H120, Live
30903.00 50% Embryo Infective Dose / 1.00 50% Embryo Infective Dose

**Pharmaceutical form:**
Lyophilisate for suspension

**Withdrawal period by route of administration:**
**Nebulisation use:**

- **Chicken (hen)**
  - All relevant tissues. 0 day

**In drinking water use:**

- **Chicken (hen)**
  - All relevant tissues. 0 day

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QI01AD06
QI01AD07

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

**Authorisation status:**
Valid

**Authorised in:**
Poland

**Package description:**
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Well-established use application (Article 13a of Directive No 2001/82/EC)

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

**Marketing authorisation date:**
4/12/2001

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

**Responsible authority:**
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

**Authorisation number:**
1232

**Date of authorisation status change:**
4/12/2001

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

## Documents

Package Leaflet

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Labelling

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

Summary of Product Characteristics

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