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AviPro THYMOVAC Lyophilisate for use in drinking water

Authorised

- Chicken anaemia virus, strain CUX-1, Live

Product identification

Medicine name:
AviPro THYMOVAC Lyophilisate for use in drinking water

Active substance:
Chicken anaemia virus, strain CUX-1, Live

Target species:
Chicken

Route of administration:
In drinking water use

Product details

Active substance and strength:
Chicken anaemia virus, strain CUX-1, Live
31622.00 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Norway

Available in:

Norway

Package description:

(ID10) 100000 Dose: Box with 10 Vial (Glass type I) each with 10000 Dose, closed with Stopfen (rubber) and Kappe (aluminium)
(ID9) 10000 Dose: Box with 1 Vial (Glass type I) with 10000 Dose, closed with Stopper (rubber) and Kappe (aluminium)
(ID8) 50000 Dose: Box with 10 Vial (Glass type I) each with 5000 Dose, closed with Stopfen (rubber) and Kappe (aluminium)
(ID7) 5000 Dose: Box with 1 Vial (Glass type I) with 5000 Dose, closed with Stopfen (rubber) and Kappe (aluminium)
(ID6) 25000 Dose: Box with 10 Vial (Glass type I) each with 2500 Dose, closed with Stopfen (rubber) and Kappe (aluminium)
(ID5) 2500 Dose: Box with 1 Vial (Glass type I) with 2500 Dose, closed with Stopfen (rubber) and Kappe (aluminium)
(ID1) 500 Dose: Box with 1 Vial (Glass type I) with 500 Dose, closed with Stopfen (rubber) and Kappe (aluminium)

(ID2) 5000 Dose: Box with 10 Vial (Glass type I) each with 500 Dose, closed with Stopfen (rubber) and Kappe (aluminium)
(ID4) 10000 Dose: Box with 10 Vial (Glass type I) each with 1000 Dose, closed with Stopfen (rubber) and Kappe (aluminium)
(ID3) 1000 Dose: Box with 1 Vial (Glass type I) with 1000 Dose, closed with Stopfen (rubber) and Kappe (aluminium)

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:
Lohmann Animal Health GmbH

Marketing authorisation date:
17/02/2009

Manufacturing sites for batch release:
Lohmann Animal Health GmbH

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
08-5862

Date of authorisation status change:
30/07/2013

Reference member state:
Germany

Procedure number:
DE/V/0247/001

Concerned member states:

Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland France Greece Hungary Iceland Ireland Italy Latvia Lithuania Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden

United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Combined File of all Documents