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
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**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Egg. no withdrawal period withdrawal period is 0 days

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:

United Kingdom (Northern Ireland)

Available in:

United Kingdom (Northern Ireland)

Package description:

(ID10): 1 Box with 10 Bottle (Glass) with 10000 Dose (100000 Dose)
(ID9): 1 Box with 1 Bottle (Glass) with 10000 Dose (10000 Dose)
(ID8): 1 Box with 10 Bottle (Glass) with 5000 Dose (50000 Dose)
(ID7): 1 Box with 1 Bottle (Glass) with 5000 Dose (5000 Dose)
(ID6): 1 Box with 10 Bottle (Glass) with 2500 Dose (25000 Dose)
(ID5): 1 Box with 1 Bottle (Glass) with 2500 Dose (2500 Dose)
(ID1): 1 Box with 1 Bottle (Glass) with 500 Dose (500 Dose)
(ID2): 1 Box with 10 Bottle (Glass) with 500 Dose (5000 Dose)
(ID4): 1 Box with 10 Bottle (Glass) with 1000 Dose (10000 Dose)
(ID3): 1 Box with 1 Bottle (Glass) with 1000 Dose (1000 Dose)

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:
Elanco GmbH

Marketing authorisation date:
28/04/2009

Manufacturing sites for batch release:
Lohmann Animal Health GmbH

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 52127/3006

Date of authorisation status change:
17/08/2021

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

Combined File of all Documents