AviPro GUMBORO VAC Lyophilisate for suspension for chicken

Not authorised

- Infectious bursal disease virus, strain Cu-1 M, Live

Product identification

Medicine name:
AviPro GUMBORO VAC Lyophilisate for suspension for chicken
Avipro Gumboro Vac Lyofilisaat voor gebruik in drinkwater
Avipro Gumboro Vac Lyophilisat pour administration dans l'eau de boisson
Avipro Gumboro Vac Lyophilisat zum Eingeben über das Trinkwasser

Active substance:
Infectious bursal disease virus, strain Cu-1 M, Live

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

Route of administration:
In drinking water use

Product details

Active substance and strength:
Infectious bursal disease virus, strain Cu-1 M, Live
5011.87 50% Embryo Infective Dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate for suspension

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

- **Chicken (broiler)**
  - Meat and offal. 0 day
- **Chicken**
  - Egg. 0 day
  - Meat and offal. 0 day
- **Chicken (for reproduction)**
  - Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD09

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

Authorisation status:
Surrendered

Authorised in:
Belgium

Package description:

(ID1): 1 Box with 1 Bottle (Glass) with 1000 Dose (1000 Dose)
(ID4): 1 Box with 10 Bottle (Glass) with 2500 Dose (25000 Dose)
(ID2): 1 Box with 10 Bottle (Glass) with 1000 Dose (10000 Dose)
(ID3): 1 Box with 1 Bottle (Glass) with 2500 Dose (2500 Dose)
(ID7): 1 Box with 1 Bottle (Glass) with 10000 Dose (10000 Dose)
(ID8): 1 Box with 10 Bottle (Glass) with 10000 Dose (100000 Dose)
(ID6): 1 Box with 10 Bottle (Glass) with 5000 Dose (50000 Dose)
(ID5): 1 Box with 1 Bottle (Glass) with 5000 Dose (5000 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:
26/06/2000

Manufacturing sites for batch release:
Lohmann Animal Health GmbH

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V214952

Date of authorisation status change:
12/05/2023

Reference member state:
Germany

Procedure number:
DE/V/0206/001

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 28/01/2022
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Package Leaflet

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