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

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 use in drinking water

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:
Revoked

Authorised in:
Portugal

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:
18/11/1999

Manufacturing sites for batch release:
Lohmann Animal Health GmbH

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
645/99 DGV

Date of authorisation status change:
1/11/2024

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

Combined File of all Documents

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