

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:

Surrendered

Authorised in:

Germany

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:

Lohmann Animal Health GmbH

Marketing authorisation date:

27/02/1998

Manufacturing sites for batch release:

Lohmann Animal Health GmbH

Responsible authority:

Paul-Ehrlich-Institut

Authorisation number:

87a/78

Date of authorisation status change:

6/06/2025

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)

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