AviPro IBD Xtreme Lyophilisate for use in drinking water for chickens

Authorised

- Infectious bursal disease virus, strain V217 (intermediate plus), Live

Product identification

Medicine name:
AviPro IBD Xtreme Lyophilisate for use in drinking water for chickens

Active substance:
Infectious bursal disease virus, strain V217 (intermediate plus), Live

Target species:
Chicken

Route of administration:
In drinking water use

Product details

Active substance and strength:
Infectious bursal disease virus, strain V217 (intermediate plus), Live
1000.00 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**
  - Meat and offal. 0 day
  - Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD09

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

France

Package description:

(ID9): 1 Box with 10 Bottle (Glass) with 5000 Dose (50000 Dose)
(ID5): 1 Box with 1 Bottle (Glass) with 10000 Dose (10000 Dose)
(ID10): 1 Box with 10 Bottle (Glass) with 10000 Dose (100000 Dose)
(ID2): 1 Box with 1 Bottle (Glass) with 1000 Dose (1000 Dose)
(ID8): 1 Box with 10 Bottle (Glass) with 2500 Dose (25000 Dose)
(ID1): 1 Box with 1 Bottle (Glass) with 500 Dose (500 Dose)
(ID6): 1 Box with 10 Bottle (Glass) with 500 Dose (5000 Dose)
(ID4): 1 Box with 1 Bottle (Glass) with 5000 Dose (5000 Dose)
(ID7): 1 Box with 10 Bottle (Glass) with 1000 Dose (10000 Dose)
(ID3): 1 Box with 1 Bottle (Glass) with 2500 Dose (2500 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:
4/09/2025

Manufacturing sites for batch release:
Lohmann Animal Health GmbH

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/6510883 3/2025

Date of authorisation status change:
4/09/2025

Reference member state:
Germany

Procedure number:
DE/V/0238/001

Concerned member states:
Bulgaria Cyprus Denmark France Greece Hungary Italy Lithuania Netherlands Poland Portugal Romania Spain

United Kingdom (Northern Ireland)

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

Documents

Package Leaflet and Labelling

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

Summary of Product Characteristics

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

Combined File of all Documents