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
AviPro IBD X Liofilizado para administração na água de bebida para galinhas

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

Authorisation status:
Valid

Authorised in:
Portugal

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:
3/04/2008

Manufacturing sites for batch release:
Lohmann Animal Health GmbH

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
798/08DIVPT

Date of authorisation status change:
1/12/2023

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

Combined File of all Documents

English (PDF)
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