Poulvac Bursa Plus Lyophilisate for suspension in drinking water

Authorised

- Infectious bursal disease virus, strain V877, Live

Product identification

Medicine name:
Poulvac Bursa Plus Lyophilisate for suspension in drinking water
Poulvac Bursa plus liofilizado para suspensão na água de bebida

Active substance:
Infectious bursal disease virus, strain V877, Live

Target species:
Chicken

Route of administration:
In drinking water use

Product details

Active substance and strength:
Infectious bursal disease virus, strain V877, Live
158.00 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**

  - Egg. 0 day

  - Meat and offal. 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:
(ID6): 1 Box with 10 Bottle (Glass) with 5000 Dose (50000 Dose)
(ID5): 1 Box with 10 Bottle (Glass) with 2000 Dose (20000 Dose)
(ID4): 1 Box with 10 Bottle (Glass) with 1000 Dose (10000 Dose)
(ID3): 1 Box with 1 Bottle (Glass) with 5000 Dose (5000 Dose)
(ID2): 1 Box with 1 Bottle (Glass) with 2000 Dose (2000 Dose)
(ID1): 1 Box with 1 Bottle (Glass) with 1000 Dose (1000 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:
Zoetis Portugal Lda.

Marketing authorisation date:

1/12/2010

Manufacturing sites for batch release:
Zoetis Manufacturing & Research Spain S.L.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
835/10 RIVPT

Date of authorisation status change:
1/12/2010

Reference member state:
Germany

Procedure number:
DE/V/0278/001

Concerned member states:
Belgium Greece Ireland Italy Lithuania Netherlands Poland Portugal Romania Slovenia United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 9/08/2022
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