

# Poulvac Bursa Plus Lyophilisate for suspension in drinking water

Authorised

- Infectious bursal disease virus, strain V877 (intermediate plus), Live

## Product identification

**Medicine name:**

Poulvac Bursa Plus Lyophilisate for suspension in drinking water

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**Active substance:**

Infectious bursal disease virus, strain V877 (intermediate plus), Live

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**Target species:**

Chicken

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**Route of administration:**

In drinking water use

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## Product details

**Active substance and strength:**

Infectious bursal disease virus, strain V877 (intermediate plus), Live  
158.00 50% Embryo Infective Dose / 1.00 Dose

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**Pharmaceutical form:**

Lyophilisate for ocular/nasal suspension/use in drinking water

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**Withdrawal period by route of administration:****In drinking water use:**

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**Chicken**

- Egg. 0 day

- Meat and offal. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01AD09

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Germany

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**Available in:**

Germany

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**Package description:**

(ID1) 1000 Dose: Box (cardboard) with 1 Bottle (Glass type I) with 1000 Dose, closed with Stopper (butyl rubber) and Kappe (aluminium)

(ID2) 2000 Dose: Box (cardboard) with 1 Bottle (Glass type I) with 2000 Dose, closed with Stopper (butyl rubber) and Kappe (aluminium)

(ID3) 5000 Dose: Box (cardboard) with 1 Bottle (Glass type I) with 5000 Dose, closed with Stopper (butyl rubber) and Kappe (aluminium)

(ID4) 10000 Dose: Box (cardboard) with 10 Bottle (Glass type I) each with 1000 Dose, closed with Stopper (butyl rubber) and Kappe (aluminium)

(ID5) 20000 Dose: Box (cardboard) with 10 Bottle (Glass type I) each with 2000 Dose, closed with Stopper (butyl rubber) and Kappe (aluminium)

(ID6) 50000 Dose: Box (cardboard) with 10 Bottle (Glass type I) each with 5000 Dose, closed with Stopper (butyl rubber) and Kappe (aluminium)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Zoetis Deutschland GmbH

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**Marketing authorisation date:**

29/07/2010

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**Manufacturing sites for batch release:**

Zoetis Manufacturing & Research Spain S.L.

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**Responsible authority:**

Paul-Ehrlich-Institut

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**Authorisation number:**

PEI.V.04760.01.1

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**Date of authorisation status change:**

22/04/2013

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0278/001

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**Concerned member states:**

Belgium Greece Ireland Italy Lithuania Netherlands Poland Portugal  
Romania Slovenia United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

Published on: 26/03/2026

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