

# Avishield IBD plus, lyophilisate for use in drinking water, for chickens

Authorised

- Infectious bursal disease virus, strain G6 (intermediate plus), Live

## Product identification

**Medicine name:**

Avishield IBD plus, lyophilisate for use in drinking water, for chickens

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**Active substance:**

Infectious bursal disease virus, strain G6 (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 G6 (intermediate plus), Live  
1.90 unit(s) / 1.00 Dose

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**Pharmaceutical form:**

Lyophilisate for use in drinking water

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

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**Chicken**

- All relevant tissues. 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:**

Spain

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**Package description:**

Cardboard box with 10 vials of 1,000 doses of vaccine. Colourless glass vials (type I) filled into 4 ml (1000 doses), which are closed with bromobutyl rubber stoppers and sealed with aluminium caps.

Cardboard box with 10 vials of 2,500 doses of vaccine. Colourless glass vials (type I) filled into 10 ml (2500 doses), which are closed with bromobutyl rubber stoppers and sealed with aluminium caps.

Cardboard box with 10 vials of 5,000 doses of vaccine. Colourless glass vials (type I) filled into 10 ml (5000 doses), which are closed with bromobutyl rubber stoppers and sealed with aluminium caps.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Izo S.r.l.

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**Marketing authorisation date:**

18/09/2020

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**Manufacturing sites for batch release:**

Genera d.d.

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

3939 ESP

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**Date of authorisation status change:**

19/09/2020

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0311/001

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**Concerned member states:**

Austria Belgium Croatia Czechia Estonia France Germany Greece Hungary  
Ireland Italy Latvia Lithuania Poland Portugal Romania Slovakia Slovenia  
Spain 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

### Summary of Product Characteristics

English (PDF)

Published on: 27/06/2024

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### Package Leaflet

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### Labelling

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