

# Avishield IB H120, lyophilisate for ocular nasal suspension/use in drinking water, for chickens

Authorised

- Infectious bronchitis virus, type Massachusetts, strain H120, Live

## Product identification

### Medicine name:

Avishield IB H120, lyophilisate for ocular nasal suspension/use in drinking water, for chickens

### Active substance:

Infectious bronchitis virus, type Massachusetts, strain H120, Live

### Target species:

Chicken

### Route of administration:

Ocular nasal use  
In drinking water use

## Product details

### Active substance and strength:

Infectious bronchitis virus, type Massachusetts, strain H120, Live

3.50 log 10 50% embryo infective dose / 1.00 Dose

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**Pharmaceutical form:**

Lyophilisate for ocularnasal suspension/use in drinking water

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**Withdrawal period by route of administration:**

**Ocularnasal use:**

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**Chicken**

- Meat and offal. 0 day

**In drinking water use:**

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**Chicken**

- Meat and offal. 0 day

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

QI01AD07

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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:**

Belgium

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**Available in:**

Belgium

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**Package description:**

Carton box with 10 vials of 5000 doses of vaccine. Colourless glass vials (type I), which are closed with bromobutyl rubber stoppers and sealed with aluminium caps. Carton box with 10 vials of 2500 doses of vaccine. Colourless glass vials (type I), which are closed with bromobutyl rubber stoppers and sealed with aluminium caps. Carton box with 10 vials of 1000 doses of vaccine. Colourless glass vials (type I), 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:**

23/02/2018

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**Manufacturing sites for batch release:**

Genera d.d.

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**Responsible authority:**

Federal Agency For Medicines And Health Products

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**Authorisation number:**

BE-V526000

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**Date of authorisation status change:**

23/02/2018

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0292/001

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**Concerned member states:**

Austria Belgium Croatia Czechia Denmark 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

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

### Package Leaflet

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

### Labelling

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

### Combined File of all Documents