Avishield IBD INT, lyophilisate for oculonasal suspension/use in drinking water for chickens

Authorised

- Infectious bursal disease virus, strain VMG 91 (intermediate), Live

Product identification

Medicine name:
Avishield IBD INT, lyophilisate for oculonasal suspension/use in drinking water for chickens
Avishield IBD INT λυοφιλοποιημένο υλικό για οφθαλμορινικό εναιώρημα/χρήση σε πόσιμο νερό για ορνίθια

Active substance:
Infectious bursal disease virus, strain VMG 91 (intermediate), Live

Target species:
Chicken

Route of administration:
Oculonasal use
Oral use

Product details

Active substance and strength:
Infectious bursal disease virus, strain VMG 91 (intermediate), Live
4.00 log10 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate for oculonasal suspension/use in drinking water

Withdrawal period by route of administration:

Oculonasal use:

- **Chicken**
  - All relevant tissues. no withdrawal period 0 days

Oral use:

- **Chicken**
  - All relevant tissues. no withdrawal period 0 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD09

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Greece

Package description:
Carton box with 10 vials of 5000 doses of vaccine
Carton box with 10 vials of 2500 doses of vaccine
Carton box with 10 vials of 1000 doses of vaccine

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Genera d.d.

Marketing authorisation date:
27/01/2019

Manufacturing sites for batch release:
Genera d.d.

Responsible authority:
National Organization For Medicines

Authorisation number:
19575/19-02-2025/K-0232601

Date of authorisation status change:
18/02/2025

Reference member state:
Netherlands

Procedure number:
NL/V/0244/001

Concerned member states:
Austria Belgium Czechia Germany Greece Hungary Ireland Italy Poland Portugal Romania Slovakia Spain

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

Documents

Combined File of all 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.