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
Avishield IBD Plus vakcina A.U.V., liofilizátum itatóvízben való használatra, házityúk számára

Active substance:
Infectious bursal disease virus, strain G6 (intermediate plus), Live

Target species:
Chicken

Route of administration:
In drinking water use

Product details

Active substance and strength:
Infectious bursal disease virus, strain G6 (intermediate plus), Live
1.90 unit(s) / 1.00 Dose

Pharmaceutical form:
Lyophilisate for use in drinking water

Withdrawal period by route of administration:
In drinking water 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:
Hungary

Package description:
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.
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 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.

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:
24/07/2020

Manufacturing sites for batch release:
Genera d.d.

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
4172/X/20 NÉBIH ÁTI

Date of authorisation status change:
24/07/2020

Reference member state:
Netherlands

Procedure number:
NL/V/0311/001

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)

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

Documents

Summary of Product Characteristics