Avishield ND +IB H120

Authorised

- Newcastle disease virus, strain La Sota, Live
- Avian infectious bronchitis virus, type Massachusetts, strain H120, Live

Product identification

Medicine name:
Avishield ND +IB H120

Active substance:
Newcastle disease virus, strain La Sota, Live

Avian infectious bronchitis virus, type Massachusetts, strain H120, Live

Target species:
Chicken

Route of administration:
Oculonasal use
In drinking water use

Product details

Active substance and strength:
Newcastle disease virus, strain La Sota, Live
6.00 log10 50% tissue culture infectious dose / 1.00 Vial

Avian infectious bronchitis virus, type Massachusetts, strain H120, Live

3.50 log 10 50% embryo infective dose / 1.00 Vial

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

Withdrawal period by route of administration:
Oculonasal use:

- **Chicken**
  - Meat and offal. 0 day

In drinking water use:

- **Chicken**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD11

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

Authorisation status:
Valid

Authorised in:
Croatia

Package description:
The vaccine is filled into colourless glass vials (type I), which are closed with rubber stoppers (brombutyl) and sealed with aluminium caps. 10x2500 dose
The vaccine is filled into colourless glass vials (type I), which are closed with rubber stoppers (brombutyl) and sealed with aluminium caps. 10x1000 dose

Additional information

Entitlement type:

Marketing Authorisation

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

Marketing authorisation holder:
Genera d.d.

Marketing authorisation date:
11/06/2018

Manufacturing sites for batch release:
Genera d.d.

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/18-01/281

Date of authorisation status change:
11/06/2018

Reference member state:
Hungary

Procedure number:
HU/V/0130/001

Concerned member states:
Croatia Slovenia

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

Documents

Summary of Product Characteristics

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