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

Authorised

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

Product identification

Medicine name:
Avishield IB H120, lyophilisate for oculonasal suspension/use in drinking water, for chickens
Avishield IB H120, liofilizat za okulonazalno suspenzijo/ uporabo v vodi za pitje za piščance

Active substance:
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:
Avian infectious bronchitis virus, type Massachusetts, strain H120, Live
3.50 log 10 50% embryo infective dose / 1.00 Dose

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

Withdrawal period by route of administration:

Oculonasal use:

- **Chicken**
  - Meat and offal. no withdrawal period 0 days

In drinking water use:

- **Chicken**
  - Meat and offal. no withdrawal period 0 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD07

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

Authorisation status:
Valid

Authorised in:
Slovenia

Package description:
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.

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 5000 doses of vaccine. Colourless glass vials (type I), 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:
29/01/2018

Manufacturing sites for batch release:
Genera d.d.

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0603/001

Date of authorisation status change:
29/01/2018

Reference member state:
Netherlands

Procedure number:
NL/V/0292/001

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)

To consult adverse reactions on veterinary medicinal products please go to 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.