Avishield ND, Lyophilisate for Oculonasal Suspension/Use in Drinking Water, for Chickens and Turkeys

Authorised

- Newcastle disease virus, strain La Sota, Live

Product identification

Medicine name:
Avishield ND, Lyophilisate for Oculonasal Suspension/Use in Drinking Water, for Chickens and Turkeys

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

Target species:
Turkey
Chicken

Route of administration:
Oculonasal use
Oral use

Product details

Active substance and strength:
Newcastle disease virus, strain La Sota, Live
6.00 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:

- **Turkey**
  - All relevant tissues. 0 day
- **Chicken**
  - All relevant tissues. 1 day

Oral use:

- **Turkey**
  - All relevant tissues. 0 day
- **Chicken**
  - All relevant tissues. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD06

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

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Package description:
Cardboard or plastic box with 10 vials of 5000 doses of vaccine. Colourless glass vials (type I), which are closed with rubber stoppers and sealed with aluminium caps.
Cardboard or plastic box with 10 vials of 1000 doses of vaccine. Colourless glass vials (type I), which are closed with rubber stoppers and sealed with aluminium caps.
Cardboard or plastic box with 10 vials of 2500 doses of vaccine. Colourless glass vials (type I), which are closed with 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:
Izo S.r.l.

Marketing authorisation date:
26/04/2016

Manufacturing sites for batch release:
Genera d.d.

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 61812/3000

Date of authorisation status change:
18/04/2023

Reference member state:
Netherlands

Procedure number:
NL/V/0300/001

Concerned member states:
Belgium Croatia Germany Greece Hungary Poland Portugal Romania Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents