Innovax-ND-H5 (--) - Concentrate and solvent for suspension for injection

Authorised

- Turkey herpesvirus, strain HVT-ND-H5 (cell-associated), expressing fusion protein gene of Newcastle disease virus and haemagglutinin gene of Avian influenza virus subtype H5, Live

Product identification

Medicine name:
Innovax-ND-H5 (--) - Concentrate and solvent for suspension for injection

Active substance:
Turkey herpesvirus, strain HVT-ND-H5 (cell-associated), expressing fusion protein gene of Newcastle disease virus and haemagglutinin gene of Avian influenza virus subtype H5, Live

Target species:
Chicken
Chicken (embryonated eggs)

Route of administration:
In ovo
Subcutaneous use

Product details

Active substance and strength:
Turkey herpesvirus, strain HVT-ND-H5 (cell-associated), expressing fusion protein gene of Newcastle disease virus and haemagglutinin gene of Avian influenza virus subtype H5, Live

Presentation_strength:10^3.3 - 10^4.6 Reference:In-house Index:0

Pharmaceutical form:
Concentrate and solvent for suspension for injection

Withdrawal period by route of administration:
In ovo:

- **Chicken**
  - Not applicable. 0 day — Zero days
- **Chicken (embryonated eggs)**
  - Not applicable. 0 day — Zero days

Subcutaneous use:

- **Chicken**
  - Not applicable. 0 day — Zero days
- **Chicken (embryonated eggs)**
  - Not applicable. 0 day — Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:
Packaging:Concentrate: ampoule (glass), Package_size:Concentrate: 1 ampoule, Content:Concentrate: 2 ml (4000 doses)
Packaging:Concentrate: ampoule (glass), Package_size:Concentrate: 1 ampoule, Content:Concentrate: 2 ml (2000 doses)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Applications in exceptional circumstances (Article 25 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
22/05/2024

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
22/05/2024

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

Documents

Combined File of all Documents

English (PDF)
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