

Innovax-ILT (--) concentrate and solvent for suspension for injection

Authorised

- Turkey herpesvirus, strain HVT/ILT-138 (cell-associated), expressing gD and gI glycoproteins genes of Infectious laryngotracheitis virus, Live

Product identification

Medicine name:

Innovax-ILT (--) concentrate and solvent for suspension for injection

Active substance:

Turkey herpesvirus, strain HVT/ILT-138 (cell-associated), expressing gD and gI glycoproteins genes of Infectious laryngotracheitis virus, Live

Target species:

Chicken

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Turkey herpesvirus, strain HVT/ILT-138 (cell-associated), expressing gD and gI glycoproteins genes of Infectious laryngotracheitis virus, Live
Presentation_strength: $10^{3.1} - 10^{4.1}$ PFU Reference: In house Comments: per dose
Index: 0

Pharmaceutical form:

Concentrate and solvent for suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Chicken

- Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD03

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)

Available in:

Austria , Belgium , Bulgaria , Cyprus , Czechia , France , Germany , Greece , Hungary , Netherlands , Poland , Spain , United Kingdom (Northern Ireland)

Package description:

Packaging: Concentrate: ampoule (glass), Package_size: Concentrate: 1 ampoule,
Content: Concentrate: 4000 doses

Packaging: Concentrate: ampoule (glass), Package_size: Concentrate: 1 ampoule,
Content: Concentrate: 2000 doses

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

3/07/2015

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:

3/07/2015

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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