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Vectormune ND (--) Suspension and solvent for suspension for injection

Authorised

- Turkey herpesvirus, strain rHVT/ND (cell-associated), encoding fusion protein gene of Newcastle disease virus (strain D-26), Live

Product identification

Medicine name:
Vectormune ND (--) Suspension and solvent for suspension for injection

Active substance:
Turkey herpesvirus, strain rHVT/ND (cell-associated), encoding fusion protein gene of Newcastle disease virus (strain D-26), Live

Target species:
Chicken
Chicken (embryonated eggs)

Route of administration:
In ovo
Subcutaneous use

Product details

Active substance and strength:

Turkey herpesvirus, strain rHVT/ND (cell-associated), encoding fusion protein gene of Newcastle disease virus (strain D-26), Live

Presentation_strength:min 2500-max 8000 PFU Reference:HSE Index:0

Pharmaceutical form:

Suspension 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

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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 , Denmark , France , Germany , Hungary , Italy , Luxembourg , Netherlands , Poland , Portugal , Romania , Slovakia , Spain , United Kingdom (Northern Ireland)

Package description:

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

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

Packaging:Ampoule (glass), Package_size:1 ampoule, Content:1000 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:

Ceva-Phylaxia Zrt.

Marketing authorisation date:

8/09/2015

Manufacturing sites for batch release:

Ceva-Phylaxia Zrt.

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:
8/09/2015

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

Documents

Combined File of all Documents

English (PDF)
Published on: 13/03/2026
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