

# Ultifend ND IBD (--) - Concentrate and solvent for suspension for injection

Authorised

- Turkey herpesvirus, strain rHVT/ND/IBD (cell-associated), expressing fusion protein gene of Newcastle disease virus and VP2 protein gene of Infectious bursal disease virus, Live

## Product identification

**Medicine name:**

Ultifend ND IBD (--) - Concentrate and solvent for suspension for injection

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**Active substance:**

Turkey herpesvirus, strain rHVT/ND/IBD (cell-associated), expressing fusion protein gene of Newcastle disease virus and VP2 protein gene of Infectious bursal disease virus, Live

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**Target species:**

Chicken

Chicken (embryonated eggs)

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**Route of administration:**

In ovo

Subcutaneous use

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## Product details

### **Active substance and strength:**

Turkey herpesvirus, strain rHVT/ND/IBD (cell-associated), expressing fusion protein gene of Newcastle disease virus and VP2 protein gene of Infectious bursal disease virus, Live

Presentation\_strength:4000 PFU/dose - 12000 PFU/dose Index:0

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### **Pharmaceutical form:**

Concentrate and solvent for suspension for injection

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### **Withdrawal period by route of administration:**

#### **In ovo:**

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#### **Chicken**

- Not applicable. 0 day Zero days

#### **Subcutaneous use:**

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#### **Chicken**

- Not applicable. 0 day Zero days

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### **Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01AD16

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### **Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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### **Authorisation status:**

Valid

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### **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)

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**Package description:**

Packaging:Concentrate: ampoule (glass), solvent: bag (PVC), Package\_size:1 ampoule, Content:4000 doses

Packaging:Concentrate: ampoule (glass), solvent: bag (PVC), Package\_size:1 ampoule, Content:1000 doses

Packaging:Concentrate: ampoule (glass), solvent: bag (PVC), Package\_size:1 ampoule, Content:2000 doses

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Ceva-Phylaxia Zrt.

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**Marketing authorisation date:**

20/04/2021

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**Manufacturing sites for batch release:**

Ceva-Phylaxia Zrt.

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

20/04/2021

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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