

# AVIFFA RTI lyofilizát na okulonazálnu suspenziu/na podanie v pitnej vode pre kurčatá a morky

Authorised

- Turkey rhinotracheitis virus, strain VCO3, Live

## Product identification

### **Medicine name:**

AVIFFA RTI lyofilizát na okulonazálnu suspenziu/na podanie v pitnej vode pre kurčatá a morky

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### **Active substance:**

Turkey rhinotracheitis virus, strain VCO3, Live

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### **Target species:**

Chicken

Turkey

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### **Route of administration:**

In drinking water use

Oculonasal use

Nebulisation use

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## Product details

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

Turkey rhinotracheitis virus, strain VCO3, Live

4.00 log<sub>10</sub> 50% cell culture infectious dose / 50.00 microlitre(s)

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### **Pharmaceutical form:**

Lyophilisate for ocular suspension/use in drinking water

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

#### **In drinking water use:**

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##### **Chicken**

- All relevant tissues. 0 day zero days (chicken and turkey)

#### **Ocular use:**

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##### **Chicken**

- All relevant tissues. 0 day zero days (chicken and turkey)

#### **Nebulisation use:**

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##### **Turkey**

- All relevant tissues. 0 day zero days

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

QI01AD01

QI01CD01

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

This information is not available for this product.

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### **Authorisation status:**

Valid

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**Authorised in:**

Slovakia

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**Package description:**

Available only in [Slovak](#)

Available only in [Slovak](#)

Available only in [Slovak](#)

Available only in [Slovak](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis reviewed according to Acquis communautaire

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**Marketing authorisation holder:**

Boehringer Ingelheim Animal Health France

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**Marketing authorisation date:**

30/05/1997

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

97/0098/97-S

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**Date of authorisation status change:**

30/05/1997

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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

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