

# Aviffa RTI liofilizátum szuszpenzióhoz pulykák és házityúkok részére

Authorised

- Turkey rhinotracheitis virus, strain VCO3, Live

## Product identification

**Medicine name:**

Aviffa RTI liofilizátum szuszpenzióhoz pulykák és házityúkok részére

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**Active substance:**

Turkey rhinotracheitis virus, strain VCO3, Live

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**Target species:**

Turkey  
Chicken

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**Route of administration:**

In drinking water/milk use  
Intraocular use  
Nebulisation use

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## Product details

**Active substance and strength:**

Turkey rhinotracheitis virus, strain VCO3, Live

2.30 log<sub>10</sub> 50% cell culture infectious dose / 1.00 unit(s)/dose

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**Pharmaceutical form:**

Oral lyophilisate

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**Withdrawal period by route of administration:**

**In drinking water/milk use:**

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**Turkey**

- Meat and offal. 0 day

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**Chicken**

- Meat and offal. 0 day

- Meat and offal. 0 day

**Intraocular use:**

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**Turkey**

- Meat and offal. 0 day

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**Chicken**

- Meat and offal. 0 day

- Meat and offal. 0 day

**Nebulisation use:**

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**Turkey**

- Meat and offal. 0 day

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

QI01AD01

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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:**

Hungary

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**Available in:**

Hungary

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**Package description:**

Available only in [Hungarian](#)

Available only in [Hungarian](#)

Available only in [Hungarian](#)

Available only in [Hungarian](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Boehringer Ingelheim Animal Health France

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**Marketing authorisation date:**

14/06/2001

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France

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**Responsible authority:**

Directorate Of Veterinary Medicinal Products

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

14/06/2001

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