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# Nobilis TRT Lyophilisate for suspension

Not  
authorised

- Turkey rhinotracheitis virus, strain BUT1#8544, Live

## Product identification

**Medicine name:**

Nobilis TRT Lyophilisate for suspension

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**Active substance:**

Turkey rhinotracheitis virus, strain BUT1#8544, Live

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**Target species:**

Turkey

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**Route of administration:**

Nebulisation use

Intraocular use

Nasal use

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## Product details

**Active substance and strength:**

Turkey rhinotracheitis virus, strain BUT1#8544, Live

0.40 50% tissue culture infectious dose / 1.00 Dose

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**Pharmaceutical form:**

Lyophilisate and solvent for ocular suspension

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**Withdrawal period by route of administration:**

**Nebulisation use:**

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**Turkey**

- Meat and offal. 0 day

**Intraocular use:**

- 

**Turkey**

- Meat and offal. 0 day

**Nasal use:**

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**Turkey**

- Meat and offal. 0 day

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

QI01CD01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Surrendered

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**Authorised in:**

Germany

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**Package description:**

(ID4): 1 Box with (1 Bottle (Glass) with 5000 Dose and 1 Bottle (-) with 0 millilitre(s))  
(5000.0 Dose, 0.0 millilitre(s))

(ID3): 1 Box with (1 Bottle (Glass) with 2500 Dose and 1 Bottle (-) with 0 millilitre(s))  
(2500.0 Dose, 0.0 millilitre(s))

(ID2): 1 Box with (1 Bottle (Glass) with 1000 Dose and 1 Bottle (-) with 0 millilitre(s))  
(1000.0 Dose, 0.0 millilitre(s))

(ID1): 1 Box with (1 Bottle (Glass) with 500 Dose and 1 Bottle (-) with 0 millilitre(s))  
(500.0 Dose, 0.0 millilitre(s))

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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:**

Intervet Deutschland GmbH

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**Marketing authorisation date:**

16/06/1997

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**Manufacturing sites for batch release:**

Intervet International B.V.

Merck Sharp & Dohme Animal Health S.L.

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**Responsible authority:**

Paul-Ehrlich-Institut

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**Authorisation number:**

189a/95

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**Date of authorisation status change:**

24/03/2021

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0205/001

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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

Summary of Product Characteristics

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