

# NEMOVAC LYOPHILISATE FOR OCULONASAL SUSPENSION/USE IN DRINKING WATER

Authorised

- Turkey rhinotracheitis virus, strain PL21, Live

## Product identification

**Medicine name:**

NEMOVAC LYOPHILISATE FOR OCULONASAL SUSPENSION/USE IN DRINKING WATER

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**Active substance:**

Turkey rhinotracheitis virus, strain PL21, Live

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**Target species:**

Chicken (layer hen)

Chicken (for reproduction)

Chicken (broiler)

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Turkey rhinotracheitis virus, strain PL21, Live

2.30 log<sub>10</sub> 50% tissue culture infectious dose / 1.00 Dose

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**Pharmaceutical form:**

Lyophilisate for oculonasal suspension/use in drinking water

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**Withdrawal period by route of administration:****Oral use:**

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**Chicken (layer hen)**

- All relevant tissues. 0 day

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**Chicken (for reproduction)**

- All relevant tissues. 0 day

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**Chicken (broiler)**

- All relevant tissues. 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:**

France

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**Available in:**

France

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**Package description:**

Box of one 1000-dose bottle

Box of ten 5000-dose bottles

Box of one 5000-dose bottles

Box of ten 2000-dose bottles

Box of one 2000-dose bottle

Box of ten1000-dose bottle,

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Boehringer Ingelheim Animal Health France

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**Marketing authorisation date:**

16/10/2001

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/5721646 5/2001

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**Date of authorisation status change:**

16/10/2011

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**Reference member state:**

France

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**Procedure number:**

FR/V/0353/001

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**Concerned member states:**

Austria Germany Greece Ireland Netherlands Portugal Spain

United Kingdom (Northern Ireland)

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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

This document does not exist in this language (English). You can find it in another language below.

### Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.

### Combined File of all Documents

eu-puar-frv0353001-mr-rpe756-en.pdf