

## P3-RHINOPLUS

Authorised

- Turkey rhinotracheitis virus, strain VCO3, Inactivated
- Newcastle disease virus, strain Ulster 2C, Inactivated
- Avian paramyxovirus 3, strain PMV3, Inactivated

### Product identification

**Medicine name:**

P3-RHINOPLUS

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**Active substance:**

Turkey rhinotracheitis virus, strain VCO3, Inactivated

Newcastle disease virus, strain Ulster 2C, Inactivated

Avian paramyxovirus 3, strain PMV3, Inactivated

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**Target species:**

Turkey

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**Route of administration:**

Subcutaneous use

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### Product details

**Active substance and strength:**

Turkey rhinotracheitis virus, strain VCO3, Inactivated

5.00 log<sub>10</sub> 50% tissue culture infectious dose / 0.50 millilitre(s)

Newcastle disease virus, strain Ulster 2C, Inactivated

8.00 log<sub>10</sub> 50% embryo infective dose / 0.50 millilitre(s)

Avian paramyxovirus 3, strain PMV3, Inactivated

6.70 log<sub>10</sub> 50% embryo infective dose / 0.50 millilitre(s)

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**Pharmaceutical form:**

Emulsion for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

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**Turkey**

- Meat and offal. 0 day

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

QI01CA02

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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:**

Italy

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**Package description:**

Available only in [Italian](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Boehringer Ingelheim Animal Health Italia S.p.A. In Breve Boehringer Ingelheim Ah It S.p.A.

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**Marketing authorisation date:**

3/11/1997

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health Italia S.p.A. In Breve Boehringer Ingelheim Ah It S.p.A.

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

22/11/2002

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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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