

# Nobilis Coryza Inac

Authorised

- Avibacterium paragallinarum, serotype A, strain 083, Inactivated
- Avibacterium paragallinarum, serotype C, strain H18, Inactivated
- Avibacterium paragallinarum, serotype B, strain Spross, Inactivated

## Product identification

**Medicine name:**

Nobilis Coryza Inac

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**Active substance:**

Avibacterium paragallinarum, serotype A, strain 083, Inactivated

Avibacterium paragallinarum, serotype C, strain H18, Inactivated

Avibacterium paragallinarum, serotype B, strain Spross, Inactivated

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**Target species:**

Chicken

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Avibacterium paragallinarum, serotype A, strain 083, Inactivated

1.00 70% Protective Dose / 0.50 millilitre(s)

Avibacterium paragallinarum, serotype C, strain H18, Inactivated

1.00 70% Protective Dose / 0.50 millilitre(s)

Avibacterium paragallinarum, serotype B, strain Spross, Inactivated

1.00 70% Protective 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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**Chicken**

- Meat and offal. no withdrawal period nul dagen

- Egg. no withdrawal period nul dagen

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

QI01AB04

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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:**

Netherlands

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**Package description:**

Available only in Dutch

Available only in Dutch

Available only in Dutch

Available only in Dutch

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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:**

Intervet Nederland B.V.

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**Marketing authorisation date:**

16/01/1992

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 7144

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**Date of authorisation status change:**

2/11/2010

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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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