

# NOBILIS RISMAVAC+CA 126, suspensija ir skiediklis injekcinei suspensijai

Authorised

- Turkey herpesvirus, strain FC-126 (cell-associated), Live
- Marek's disease virus, serotype 1, strain CVI-988 (Rispens, cell-associated), Live

## Product identification

**Medicine name:**

NOBILIS RISMAVAC+CA 126, suspensija ir skiediklis injekcinei suspensijai

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**Active substance:**

Turkey herpesvirus, strain FC-126 (cell-associated), Live

Marek's disease virus, serotype 1, strain CVI-988 (Rispens, cell-associated), Live

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**Target species:**

Chicken

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

**Active substance and strength:**

Turkey herpesvirus, strain FC-126 (cell-associated), Live

3.00 log<sub>10</sub> plaque forming unit(s) / 1.00 Dose

Marek's disease virus, serotype 1, strain CVI-988 (Rispens, cell-associated), Live

3.00 log<sub>10</sub> plaque forming unit(s) / 1.00 Dose

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**Pharmaceutical form:**

Suspension and solvent for suspension for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Chicken**

- Meat and offal. 0 day

**Subcutaneous use:**

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**Chicken**

- Meat and offal. 0 day

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

QI01AD03

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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:**

Lithuania

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**Package description:**

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis reviewed according to Acquis communautaire

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

12/11/2002

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**Manufacturing sites for batch release:**

INTERVET INTERNATIONAL B.V.

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**Responsible authority:**

State Food And Veterinary Service

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

27/10/2025

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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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