

# CRYOMAREX RISPENS + HVT

## injekčná suspenzia pre kurčatá

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:**

CRYOMAREX RISPENS + HVT injekčná suspenzia pre kurčatá

**Active substance:**

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

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

**Target species:**

Chicken

**Route of administration:**

Intramuscular use

Subcutaneous use

## Product details

**Active substance and strength:**

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

3.00 log<sub>10</sub> plaque forming unit(s) / 0.20 millilitre(s)

Marek's disease virus, serotype 1, strain CVI-988 (Rispens, cell-associated), Live  
3.00 log<sub>10</sub> plaque forming unit(s) / 0.20 millilitre(s)

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Chicken**

- All relevant tissues. 0 day zero days

**Subcutaneous use:**

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**Chicken**

- All relevant tissues. 0 day zero days

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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:**

Slovakia

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**Package description:**

Available only in [Slovak](#)

Available only in [Slovak](#)

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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 France

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**Marketing authorisation date:**

13/05/1999

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

97/035/99-S

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**Date of authorisation status change:**

13/05/1999

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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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