

# Cevac MD Rispens concentrate and solvent for suspension for injection for chickens

Authorised

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

## Product identification

**Medicine name:**

Cevac MD Rispens concentrate and solvent for suspension for injection for chickens

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**Active substance:**

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:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Marek's disease virus, serotype 1, strain CVI-988 (Rispens, cell-associated), Live  
5000.00 plaque forming unit / 1.00 Dose

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**Pharmaceutical form:**

Concentrate and solvent for suspension for injection

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

Sweden

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**Package description:**

bag containing 1600 ml of solvent

bag containing 1200 ml of solvent

bag containing 1000 ml of solvent

bag containing 800 ml of solvent

bag containing 400 ml of solvent

bag containing 200 ml of solvent

1 ampoule containing 4000 doses

1 ampoule containing 2000 doses

1 ampoule containing 1000 doses

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ceva-Phylaxia Zrt.

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**Marketing authorisation date:**

22/01/2021

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**Manufacturing sites for batch release:**

Ceva-Phylaxia Zrt.

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**Responsible authority:**

Swedish Medical Products Agency

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**Authorisation number:**

57897

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**Date of authorisation status change:**

22/01/2021

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0312/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
France Germany Greece Hungary Ireland Italy Latvia Lithuania Netherlands  
Poland Portugal Romania Slovakia Slovenia Sweden  
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

### Package Leaflet

English (PDF)

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### Summary of Product Characteristics

English (PDF)

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### Labelling

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### Combined File of all Documents