

# RUVAX, süstesuspension

Not authorised

- *Erysipelothrix rhusiopathiae*, serotype 2, Inactivated

## Product identification

**Medicine name:**

RUVAX, süstesuspension

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**Active substance:**

*Erysipelothrix rhusiopathiae*, serotype 2, Inactivated

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**Target species:**

Pig

Sheep

Turkey

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

**Active substance and strength:**

*Erysipelothrix rhusiopathiae*, serotype 2, Inactivated

1.00 enzyme-linked immunosorbent assay unit / 1.00 dose

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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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**Pig**

- Meat and offal. 0 day

**Subcutaneous use:**

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**Pig**

- Meat and offal. 0 day

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**Sheep**

- Meat and offal, milk. 0 day

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**Turkey**

- Meat and offal. 0 day

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

QI09AB03

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Surrendered

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**Authorised in:**

Estonia

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**Package description:**

Available only in Estonian

Available only in Estonian

Available only in Estonian

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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:**

5/02/2004

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France SCS

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**Responsible authority:**

State Agency Of Medicines

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**Authorisation number:**

1200

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**Date of authorisation status change:**

27/04/2023

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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

Summary of Product Characteristics

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