

RABISIN,  $\geq 2,09 \log_{10}$  OD50 i  $\geq 1$   
I.J., suspenzija za injekciju za pse,  
mačke, pitome vretice, konje,  
govedo i ovce

Authorised

- Rabies virus, strain G52, Inactivated

## Product identification

### Medicine name:

RABISIN,  $\geq 2,09 \log_{10}$  OD50 i  $\geq 1$  I.J., suspenzija za injekciju za pse, mačke, pitome vretice, konje, govedo i ovce

### Active substance:

Rabies virus, strain G52, Inactivated

### Target species:

Dog  
Cat  
Ferret  
Cattle  
Sheep  
Horse

### Route of administration:

Subcutaneous use  
Intramuscular use

## Product details

### **Active substance and strength:**

Rabies virus, strain G52, Inactivated

2.09 international unit(s) / 1.00 millilitre(s)

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### **Pharmaceutical form:**

Suspension for injection

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

#### **Subcutaneous use:**

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##### **Cattle**

- Meat and offal. 0 day
- Milk. 0 day

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##### **Sheep**

- Meat and offal. 0 day
- Milk. 0 day

#### **Intramuscular use:**

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##### **Horse**

- Meat and offal. 0 day

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##### **Cattle**

- Meat and offal. 0 day
- Milk. 0 day

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##### **Sheep**

- Meat and offal. 0 day
  - Milk. 0 day
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI07AA02

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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:**

Croatia

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**Available in:**

Croatia

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**Package description:**

Available only in [Croatian](#)

Available only in [Croatian](#)

Available only in [Croatian](#)

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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:**

19/10/2018

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France

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**Responsible authority:**

Ministry Of Agriculture Veterinary And Food Safety Directorate

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**Authorisation number:**

UP/I-322-05/16-01/490

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**Date of authorisation status change:**

17/01/2025

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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

Summary of Product Characteristics

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