

# Versiguard Rabies, Suspension for injection

Authorised

- Rabies virus, strain SAD Vnukovo-32, Inactivated

## Product identification

**Medicine name:**

Versiguard Rabies, Suspension for injection

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**Active substance:**

Rabies virus, strain SAD Vnukovo-32, Inactivated

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**Target species:**

Cat

Cattle

Pig

Sheep

Goat

Horse

Ferret

Dog

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

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

Rabies virus, strain SAD Vnukovo-32, Inactivated  
2.00 international unit(s) / 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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#### **Cattle**

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

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#### **Pig**

- Meat and offal. 0 day

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#### **Sheep**

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

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#### **Goat**

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

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#### **Horse**

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

#### **Subcutaneous use:**

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**Cattle**

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

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**Pig**

- Meat and offal. 0 day

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**Sheep**

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

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**Goat**

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

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**Horse**

- Milk. 0 hour
- Meat and offal. 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:**

Germany

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**Available in:**

Germany

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**Package description:**

Glass Vial 1 x 1.0 Dose  
Glass Vial 10 x 10.0 Dose  
Glass Vial 10 x 1.0 Dose

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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:**

Zoetis Deutschland GmbH

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**Marketing authorisation date:**

3/05/2006

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**Manufacturing sites for batch release:**

Bioveta a.s.

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**Responsible authority:**

Paul-Ehrlich-Institut

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**Authorisation number:**

PEI.V.03351.01.1

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**Date of authorisation status change:**

31/01/2011

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**Reference member state:**

Czechia

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**Procedure number:**

CZ/V/0100/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Denmark Estonia Finland France

Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg  
Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain  
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

Combined File of all Documents

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Labelling

2603624-spcde-20210301.pdf