

# 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:**

France

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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 France

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**Marketing authorisation date:**

9/05/2006

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**Manufacturing sites for batch release:**

Bioveta a.s.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/8347931 9/2006

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**Date of authorisation status change:**

9/05/2011

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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

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

This document does not exist in this language (English). You can find it in another language below.

### Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.

### Labelling