File downloaded on 2026-11-16
**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000053940

# Versiguard Rabies, Suspension for injection

Authorised

- Rabies virus, strain SAD Vnukovo-32, Inactivated

## Product identification

**Medicine name:**
Versiguard Rabies, Suspension for injection

**Active substance:**
Rabies virus, strain SAD Vnukovo-32, Inactivated

**Target species:**
Cat
Cattle
Pig
Sheep
Goat
Horse
Ferret
Dog

**Route of administration:**
Intramuscular use
Subcutaneous use

## Product details

**Active substance and strength:**
Rabies virus, strain SAD Vnukovo-32, Inactivated
2.00 international unit(s) / 1.00 Dose

---

**Pharmaceutical form:**
Suspension for injection

---

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

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Sheep**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Goat**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Horse**
  - Milk. 0 hour
  - Meat and offal. 0 day

**Subcutaneous use:**

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Sheep**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Goat**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Horse**
  - Milk. 0 hour
  - Meat and offal. 0 day

---

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QI07AA02

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Norway

**Available in:**
Norway

**Package description:**
Glass Vial 1 x 1.0 Dose
Glass Vial 10 x 10.0 Dose
Glass Vial 10 x 1.0 Dose

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Full application (Article 12(3) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Zoetis Animal Health ApS

**Marketing authorisation date:**
22/05/2009

**Manufacturing sites for batch release:**
Bioveta a.s.

**Responsible authority:**
Norwegian Medical Products Agency

**Authorisation number:**
05-3851

**Date of authorisation status change:**
29/03/2011

**Reference member state:**
Czechia

**Procedure number:**
CZ/V/0100/001

**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)

---

To consult adverse reactions on veterinary medicinal products please go to 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

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