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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:
Czechia

Available in:
Czechia

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 Ceska Republika s.r.o.

Marketing authorisation date:
2/11/2005

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/040/05-C

Date of authorisation status change:
2/11/2005

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

English (PDF)
Published on: 11/09/2023
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Package Leaflet

English (PDF)
Published on: 11/09/2023
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Labelling

English (PDF)
Published on: 14/03/2026
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