Versiguard Rabies, Suspension for injection

Authorised

- Rabies virus, strain SAD Vnukovo-32, Inactivated

Product identification

Medicine name:
Versiguard Rabies, Suspension for injection
Versiguard Rabies suspensija injekcijām

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:

- **Cat**
- **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
- **Ferret**

Subcutaneous use:

- **Dog**
- **Cat**
- **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
- **Ferret**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AA02

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

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Authorisation status:
Valid

Authorised in:
Latvia

Package description:
Available only in Latvian
Available only in Latvian
Available only in Latvian

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 Belgium

Marketing authorisation date:
26/03/2006

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/MRP/06/1668

Date of authorisation status change:
26/03/2006

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

Combined File of all Documents

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