

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:

United Kingdom (Northern Ireland)

Available in:

United Kingdom (Northern Ireland)

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 Belgium S.A.

Marketing authorisation date:

3/12/2015

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 60021/3039

Date of authorisation status change:

13/06/2024

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

Labelling