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# Biocan R

Authorised

- Rabies virus, strain SAD Vnukovo-32, Inactivated

## Product identification

**Medicine name:**

Biocan R

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**Active substance:**

Rabies virus, strain SAD Vnukovo-32, Inactivated

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**Target species:**

Dog

Cat

Cattle

Pig

Sheep

Goat

Horse

Ferret

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**Route of administration:**

Subcutaneous use

Intramuscular use

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## Product details

**Active substance and strength:**

Rabies virus, strain SAD Vnukovo-32, Inactivated  
2.00 international unit(s)/millilitre / 1.00 Dose

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 0 day

•

**Pig**

- Meat and offal. 0 day

•

**Sheep**

- Meat and offal. 0 day

•

**Goat**

- Meat and offal. 0 day

•

**Horse**

- Meat and offal. 0 day

**Intramuscular use:**

•

**Cattle**

- Meat and offal. 0 day

•

**Pig**

- Meat and offal. 0 day

•

**Sheep**

- Meat and offal. 0 day

•

**Goat**

- Meat and offal. 0 day

•

**Horse**

- 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:**

Cyprus

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**Package description:**

The vaccine is supplied in Type I glass vials complying with Ph Eur, sealed with a rubber stopper and aluminium cap.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Bioveta a.s.

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**Marketing authorisation date:**

9/03/2010

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**Manufacturing sites for batch release:**

Bioveta a.s.

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**Responsible authority:**

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

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**Authorisation number:**

CY00198V

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**Date of authorisation status change:**

4/12/2013

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**Reference member state:**

Hungary

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**Procedure number:**

HU/V/0107/001

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**Concerned member states:**

Cyprus Estonia Slovenia

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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

English (PDF)

Published on: 21/12/2021

Updated on: 13/03/2026

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