

Rabadrop, Oral suspension

Authorised

- Rabies virus, strain SAD, Live

Product identification

Medicine name:

Rabadrop, Oral suspension

Active substance:

Rabies virus, strain SAD, Live

Target species:

Fox

Raccoon dog

Route of administration:

Oral use

Product details

Active substance and strength:

Rabies virus, strain SAD, Live

8.50 log₁₀ 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:

Oral suspension

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07BD

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Package description:

Plastic Sachet 2 x 350.0 Dose

Plastic Sachet 1 x 700.0 Dose

Plastic Sachet 1 x 30.0 Dose

Paper Box 30 x 20.0 Dose

Paper Box 1 x 20.0 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:

Bioveta a.s.

Marketing authorisation date:

9/10/2019

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

National Organization For Medicines

Authorisation number:

17588/17-02-2025/K-0235801

Date of authorisation status change:

16/02/2025

Reference member state:

Czechia

Procedure number:

CZ/V/0149/001

Concerned member states:

Bulgaria Croatia Estonia Finland Germany Greece Hungary Latvia Lithuania
Poland Romania Slovakia Slovenia

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)

Published on: 14/03/2026

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Package Leaflet

English (PDF)

Published on: 14/03/2026

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Labelling

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