File downloaded on 2026-11-02
Source URL: https://medicines.health.europa.eu/veterinary/en/600000054539

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:
Finland

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:
1/03/2020

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Finnish Medicines Agency

Authorisation number:
36128

Date of authorisation status change:
1/03/2020

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

eu-puar-czv0149001-mr-rabadrop-en.pdf