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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:
Lithuania

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:
23/10/2019

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/19/2555/001-005

Date of authorisation status change:
10/06/2021

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

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

Package Leaflet

Labelling

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