

Rabadrop, Oral suspension

Authorised

- Rabies virus, strain SAD, Live

Product identification

Medicine name:

Rabadrop, Oral suspension

RABADROP suspensija iekšķīgai lietošanai

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 tissue culture infective dose 50 / 1.00 Dose

Pharmaceutical form:

Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Fox**
- **Raccoon dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07BD

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Latvia

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:

3/10/2019

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/DCP/19/0059

Date of authorisation status change:

3/10/2019

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

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Package Leaflet

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Labelling

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