Rabadrop, Oral suspension

Authorised

- Rabies virus, strain SAD, Live

Product identification

Medicine name:
Rabadrop, Oral suspension
RABADROP, перорална суспензия

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:
Bulgaria

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:
13/08/2019

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Bulgarian Agency For Food Safety

Authorisation number:
0022-2908

Date of authorisation status change:
13/08/2019

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

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

Package Leaflet and Labelling

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