Rabadrop, Oral suspension

Authorised

- Rabies virus, strain SAD, Live

Product identification

Medicine name:
Rabadrop, Oral suspension
RABADROP, perorálna suspenzia

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:
Slovakia

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:
18/11/2019

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/031/DC/19-S

Date of authorisation status change:
22/07/2022

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

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