Source URL: https://medicines.health.europa.eu/veterinary/en/600000054527

Rabadrop, Oral suspension

Authorised

• Rabies virus, strain SAD, Live

Product identification

Medicine name:

Rabadrop, Oral suspension RABADROP, suspensie orală

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

Withdrawal period by route of administration: Oral use: Fox Raccoon dog Anatomical therapeutic chemical veterinary (ATCvet) codes: OI07BD Legal status of supply: Veterinary medicinal product subject to veterinary prescription **Authorisation status:** Valid **Authorised in:** Romania 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:

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

240133

Date of authorisation status change:

15/10/2024

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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