

# Rabitec (-- ) - Oral suspension

Authorised

- Rabies virus, strain SPBN GASGAS, Live

## Product identification

**Medicine name:**

Rabitec (-- ) - Oral suspension

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**Active substance:**

Rabies virus, strain SPBN GASGAS, Live

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**Target species:**

Fox

Raccoon dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Rabies virus, strain SPBN GASGAS, Live

Presentation\_strength:10<sup>6.8</sup> to 10<sup>8.1</sup> FFU Reference:Hse Index:0

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**Pharmaceutical form:**

Oral suspension

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI07BD

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

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**Package description:**

Packaging:4 × 200 baits; suspension: blister (PVC-Alu); bait: matrix,

Package\_size:800 baits, Content:Suspension: 1.7 ml (one dose); bait: PE bag

Packaging:800 baits; suspension: blister (PVC-Alu); bait: matrix, Package\_size:800 baits, Content:Suspension: 1.7 ml (one dose); bait: PE sleeve

Packaging:40 × 20 baits; suspension: blister (PVC-Alu); bait: matrix,

Package\_size:800 baits, Content:Suspension: 1.7 ml (one dose); bait: PE bag

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ceva Sante Animale

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**Marketing authorisation date:**

1/12/2017

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**Manufacturing sites for batch release:**

Ceva Tiergesundheit (Riems) GmbH

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

1/12/2017

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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