

Purevax Rabies (--) - Suspension for injection

Authorised

- Canarypox virus, strain vCP65, expressing glycoprotein G gene of Rabies virus, Live

Product identification

Medicine name:

Purevax Rabies (--) - Suspension for injection

Active substance:

Canarypox virus, strain vCP65, expressing glycoprotein G gene of Rabies virus, Live

Target species:

Cat

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Canarypox virus, strain vCP65, expressing glycoprotein G gene of Rabies virus, Live
Presentation_strength: $\geq 10^{6.8}$ FAID₅₀ Reference:Hse Index:0

Pharmaceutical form:

Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI06AX

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

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)

Available in:

Belgium , Czechia , France , Luxembourg , Poland , Slovakia , Spain

Package description:

Packaging:Vial (glass) with a butyl elastomer closure, sealed with an aluminium cap,

Package_size:2 vials, Content:1 ml

Packaging:Vial (glass) with a butyl elastomer closure, sealed with an aluminium cap.,

Package_size:50 vials, Content:1 ml

Packaging:Vial (glass) with a butyl elastomer closure, sealed with an aluminium cap.,

Package_size:10 vials, Content:1 ml

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:

Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:

18/02/2011

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

29/07/2011

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

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ema-puar-purevax-rabies-v-2003-par-en.pdf