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:≥ $10^{6,8}$ $FAID_{50}$ 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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