

Purevax FeLV (--) - Suspension for injection

Authorised

- Canarypox virus, strain vCP97, expressing env and gag genes and a portion of the pol gene of Feline leukemia virus, Live

Product identification

Medicine name:

Purevax FeLV (--) - Suspension for injection

Active substance:

Canarypox virus, strain vCP97, expressing env and gag genes and a portion of the pol gene of Feline leukemia virus, Live

Target species:

Cat

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Canarypox virus, strain vCP97, expressing env and gag genes and a portion of the pol gene of Feline leukemia virus, Live

Presentation_strength: $\geq 10^{7.2}$ CCID₅₀ Reference:Hse Index:0

Pharmaceutical form:

Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI06AD

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 , France , Luxembourg , Spain

Package description:

Packaging:Bottle (glass), Package_size:10 bottles, Content:0.5 ml

Packaging:Bottle (glass), Package_size:20 bottles, Content:0.5 ml

Packaging:Bottle (glass), Package_size:50 bottles, Content:0.5 ml

Packaging:Bottle (glass), Package_size:50 bottles, Content:1 ml

Packaging:Bottle (glass), Package_size:20 bottles, Content:1 ml

Packaging:Bottle (glass), Package_size:10 bottles, Content:1 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:

Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:

13/04/2000

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:

18/03/2021

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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