Source URL: https://medicines.health.europa.eu/veterinary/en/60000000561

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:≥10^{7,2} CCID₅₀ Reference:Hse Index:0

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

Cat

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)

Published on: 8/11/2024

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