Purevax RCP FeLV (--) - Lyophilisate and solvent for suspension for injection

Authorised

- Feline calicivirus, strains 431 and G1, Inactivated
- Feline panleucopenia virus, strain PLI IV, Live
- Felid herpesvirus 1, strain F2, Live
- 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 RCP FeLV (--) - Lyophilisate and solvent for suspension for injection

Active substance:
Feline calicivirus, strains 431 and G1, Inactivated

Feline panleucopenia virus, strain PLI IV, Live

Felid herpesvirus 1, strain F2, Live

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:

Feline calicivirus, strains 431 and G1, Inactivated

Presentation_strength:≥ 2.0 ELISA U Reference:HSE Index:0

Feline panleucopenia virus, strain PLI IV, Live

Presentation_strength:≥$10^{3,5}$ $CCID_{50}$ Reference:HSE Index:1

Felid herpesvirus 1, strain F2, Live

Presentation_strength:≥$10^{4,9}$ $CCID_{50}$ Reference:HSE Index:2

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

Presentation_strength:≥$10^{7,2}$ $CCID_{50}$ Reference:HSE Index:3

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI06AH10

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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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)

Package description:

Packaging:Lyophilisate: bottle (glass); Solvent: bottle (glass),

Package_size:Lyophilisate: 10 bottles; Solvent: 10 bottles, Content:Lyophilisate: 1

dose; Solvent: 0.5 ml
Packaging:Lyophilisate: bottle (glass); Solvent: bottle (glass),
Package_size:Lyophilisate: 50 bottles; Solvent: 50 bottles, Content:Lyophilisate: 1 dose; Solvent: 0.5 ml
Packaging:Lyophilisate: bottle (glass); Solvent: bottle (glass),
Package_size:Lyophilisate: 50 bottles; Solvent: 50 bottles, Content:Lyophilisate: 1 dose; Solvent: 1 ml
Packaging:Lyophilisate: bottle (glass); Solvent: bottle (glass),
Package_size:Lyophilisate: 10 bottles; Solvent: 10 bottles, Content:Lyophilisate: 1 dose; Solvent: 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:
23/02/2005

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:
23/02/2005

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: 10/07/2023
Updated on: 22/05/2024
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ema-puar-purevax-rcp-felv-v-089-par-en.pdf
