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Leucofeligen FeLV/RCP (--) - Lyophilisate and suspension for suspension for injection

Authorised

- Feline panleucopenia virus, strain LR 72, Live
- Feline calicivirus, strain F9, Live
- Felid herpesvirus 1, strain F2, Live
- Feline leukemia virus, envelope protein p45

Product identification

Medicine name:
Leucofeligen FeLV/RCP (--) - Lyophilisate and suspension for suspension for injection

Active substance:
Feline panleucopenia virus, strain LR 72, Live

Feline calicivirus, strain F9, Live

Felid herpesvirus 1, strain F2, Live

Feline leukemia virus, envelope protein p45

Target species:
Cat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Feline panleucopenia virus, strain LR 72, Live

Presentation_strength:$10^{3.7}$-$10^{4.5}$ $CCID_{50}$ Reference:Hse Comments:Freeze dried component Index:0

Feline calicivirus, strain F9, Live

Presentation_strength:$10^{4.6}$-$10^{6.1}$ $CCID_{50}$ Reference:Hse Comments:Freeze dried component Index:1

Felid herpesvirus 1, strain F2, Live

Presentation_strength:$10^{5.0}$-$10^{6.6}$ $CCID_{50}$ Reference:Hse Comments:Freeze dried component Index:2

Feline leukemia virus, envelope protein p45

102.00 microgram(s) / 1.00 Vial

Pharmaceutical form:

Lyophilisate and suspension for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI06AH07

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)

Available in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Poland , Portugal , Romania , Slovakia , Slovenia , United Kingdom (Northern Ireland)

Package description:
Packaging:Lyophilisate: vial (glass); Suspension: glass vial (glass), Package_size:Lyophilisate: 10 vials + Suspension: 10 vials, Content:Lyophilisate: 1 dose; Suspension: 1 ml
Packaging:Lyophilisate: vial (glass); Suspension: glass vial (glass), Package_size:Lyophilisate: 50 vials + Suspension: 50 vials, Content:Lyophilisate: 1 dose; Suspension: 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:
Virbac

Marketing authorisation date:
25/06/2009

Manufacturing sites for batch release:
Virbac

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
25/06/2009

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: 19/10/2026
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