Feligen CR/P Vivant Lyophilisat et solvant pour solution injectable

Authorised

- Felid herpesvirus 1, strain F2, Live
- Feline panleucopenia virus, strain LR 72, Live
- Feline calicivirus, strain F9, Live

Product identification

Medicine name:
Feligen CR/P Vivant Lyophilisat et solvant pour solution injectable

Active substance:
Felid herpesvirus 1, strain F2, Live

Feline panleucopenia virus, strain LR 72, Live

Feline calicivirus, strain F9, Live

Target species:
Cat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Felid herpesvirus 1, strain F2, Live

5.00 log10 50% cell culture infectious dose / 1.00 Vial

Feline panleucopenia virus, strain LR 72, Live
3.70 log10 50% cell culture infectious dose / 1.00 Vial

Feline calicivirus, strain F9, Live
4.60 log10 50% cell culture infectious dose / 1.00 Vial

Pharmaceutical form:
Lyophilisate and solvent for solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI06AD04

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
10 x 1 dose lyophilsate + 10 x 1 ml solvent. The vials is sealed with an elastomer cap and an aluminium bottle cap.
30 x 1 dose lyophilsate + 30 x 1 ml solvent. The vials is sealed with an elastomer cap and an aluminium bottle cap.
50 x 1 dose lyophilsate + 50 x 1 ml solvent. The vials is sealed with an elastomer cap and an aluminium bottle cap.
1 x 1 dose lyophilsate + 1 x 1 ml solvent. The vials is sealed with an elastomer cap and an aluminium bottle cap.

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:
1/03/1988

Manufacturing sites for batch release:
Virbac

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V140734

Date of authorisation status change:
16/05/2018

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Package Leaflet

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Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Labelling

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