

Biofel PCH, Emulsion for injection

Authorised

- Feline panleucopenia virus, strain FPV Bio 7, Inactivated
- Felid herpesvirus 1, strain FHV-1 Bio-9, Inactivated
- Feline calicivirus, strain FCV F9 Bio-8, Inactivated

Product identification

Medicine name:

Biofel PCH, Emulsion for injection

Biofel PCH injekčná emulzia pre mačky

Active substance:

Feline panleucopenia virus, strain FPV Bio 7, Inactivated

Felid herpesvirus 1, strain FHV-1 Bio-9, Inactivated

Feline calicivirus, strain FCV F9 Bio-8, Inactivated

Target species:

Cat

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Feline panleucopenia virus, strain FPV Bio 7, Inactivated

1.00 relative potency / 1.00 Dose

Felid herpesvirus 1, strain FHV-1 Bio-9, Inactivated

1.00 relative potency / 1.00 Dose

Feline calicivirus, strain FCV F9 Bio-8, Inactivated
1.00 relative potency / 1.00 Dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Cat

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI06AA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

Glass Vial 1 x 2.0 Dose

Glass Vial 1 x 10.0 Dose

Glass Vial 1 x 20.0 Dose

Glass Vial 1 x 100.0 Dose

Glass Vial 1 x 5.0 Dose

Glass Vial 5 x 5.0 Dose

Glass Vial 10 x 5.0 Dose

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:

Bioveta a.s.

Marketing authorisation date:

1/07/2019

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

97/014/MR/19-S

Date of authorisation status change:

1/07/2019

Reference member state:

Czechia

Procedure number:

CZ/V/0156/001

Concerned member states:

Slovakia

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

Documents

Combined File of all Documents

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