Biofel PCHR Emulsja do wstrzykiwań

Authorised

- Feline panleucopenia virus, strain FPV Bio 7, Inactivated
- Feline calicivirus, strain FCV F9 Bio-8, Inactivated
- Felid herpesvirus 1, strain FHV-1 Bio-9, Inactivated
- Rabies virus, strain SAD Vnukovo-32, Inactivated

Product identification

Medicine name:
Biofel PCHR Emulsja do wstrzykiwań

Active substance:
Feline panleucopenia virus, strain FPV Bio 7, Inactivated

Feline calicivirus, strain FCV F9 Bio-8, Inactivated

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

Rabies virus, strain SAD Vnukovo-32, Inactivated

Target species:
Cat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Feline panleucopenia virus, strain FPV Bio 7, Inactivated
100000.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Feline calicivirus, strain FCV F9 Bio-8, Inactivated
31622800.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Felid herpesvirus 1, strain FHV-1 Bio-9, Inactivated
10000000.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Rabies virus, strain SAD Vnukovo-32, Inactivated
1.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Emulsion for injection/infusion

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI06AA09

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Poland

Package description:

Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish

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:
Grabikowski-Grabikowska Przedsiebiorstwo Produkcyjno-Handlowo-Uslugowe Inex Sp. j.

Marketing authorisation date:
12/12/2014

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2395

Date of authorisation status change:
12/12/2014

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

Documents

Labelling

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

Summary of Product Characteristics

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

Package Leaflet

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