BIOFEL PCHR ενέσιμο γαλάκτωμα για γάτες

Authorised

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

Product identification

Medicine name:
BIOFEL PCHR ενέσιμο γαλάκτωμα για γάτες

Active substance:
Rabies virus, strain SAD Vnukovo-32, Inactivated

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

Feline calicivirus, strain FCV F9 Bio-8, Inactivated

Feline panleucopenia virus, strain FPV Bio 7, Inactivated

Target species:
Cat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

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

Felid herpesvirus 1, strain FHV-1 Bio-9, Inactivated
100000.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Feline calicivirus, strain FCV F9 Bio-8, Inactivated
316228.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Feline panleucopenia virus, strain FPV Bio 7, Inactivated
1000.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cat**
  - Not applicable. no withdrawal period

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:

Greece

Package description:

Available only in Greek

Available only in Greek

Available only in Greek
Available only in Greek
Available only in Greek
Available only in Greek
Available only in Greek

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone)

Marketing authorisation holder:
Bioveta a.s.

Marketing authorisation date:
17/05/2022

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
National Organization For Medicines

Authorisation number:
69249/19-06-2025/K-0267001

Date of authorisation status change:
18/06/2025

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

Documents

Summary of Product Characteristics

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Package Leaflet

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