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Fevaxyn Quatrifel, Injeksjonsvæske, suspensjon, til katt

Not authorised

- Feline panleucopenia virus, strain CU4, Inactivated
- Feline calicivirus, strain 255, Inactivated
- Felid herpesvirus 1, strain 605, Inactivated
- Chlamydia felis, strain Cello, Inactivated

Product identification

Medicine name:
Fevaxyn Quatrifel, Injeksjonsvæske, suspensjon, til katt

Active substance:
Feline panleucopenia virus, strain CU4, Inactivated
Feline calicivirus, strain 255, Inactivated
Felid herpesvirus 1, strain 605, Inactivated
Chlamydia felis, strain Cello, Inactivated

Target species:
Cat

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Feline panleucopenia virus, strain CU4, Inactivated
8.50 relative potency / 1.00 millilitre(s)

Feline calicivirus, strain 255, Inactivated
1.26 relative potency / 1.00 millilitre(s)

Felid herpesvirus 1, strain 605, Inactivated
1.39 relative potency / 1.00 millilitre(s)

Chlamydia felis, strain Cello, Inactivated
1.69 relative potency / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI06AL02

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

Authorisation status:
Surrendered

Authorised in:
Norway

Package description:
Available only in Norwegian
Available only in Norwegian

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:
Zoetis Animal Health ApS

Marketing authorisation date:
28/09/1995

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
8127

Date of authorisation status change:
1/10/2010

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

Documents

Summary of Product Characteristics

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