Fevaxyn Pentofel (--) - Suspension for injection

Not authorised

- Feline leukemia virus, Strain AH 927/1161E, Inactivated
- Felid herpesvirus 1, strain 605, Inactivated
- Feline calicivirus, strain 255, Inactivated
- Chlamydia felis, strain Cello, Inactivated
- Feline panleucopenia virus, strain CU4, Inactivated

Product identification

Medicine name:
Fevaxyn Pentofel (--) - Suspension for injection

Active substance:
Feline leukemia virus, Strain AH 927/1161E, Inactivated

Felid herpesvirus 1, strain 605, Inactivated

Feline calicivirus, strain 255, Inactivated

Chlamydia felis, strain Cello, Inactivated

Feline panleucopenia virus, strain CU4, Inactivated

Target species:
This information is not available for this product.

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Feline leukemia virus, Strain AH 927/1161E, Inactivated

Presentation_strength:≥ 1.45 RP Reference:Hse Index:0

Felid herpesvirus 1, strain 605, Inactivated
1.60 relative potency / 1.00 Syringe

Feline calicivirus, strain 255, Inactivated
1.65 relative potency / 1.00 Syringe

Chlamydia felis, strain Cello, Inactivated
2.00 relative potency / 1.00 Syringe

Feline panleucopenia virus, strain CU4, Inactivated
9.50 relative potency / 1.00 Syringe

Pharmaceutical form:

Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI06AL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Surrendered

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Zoetis Belgium SA

Marketing authorisation date:
5/02/1997

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
4/10/2024

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

Documents

Combined File of all Documents

English (PDF)
Published on: 24/10/2024
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