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Feliserin Plus Injektionslösung für Katzen und Hunde

Authorised

- HORSE PROTEINS
- Immunoglobulins against feline panleukopenia virus, Equine
- Immunoglobulins against Felid herpesvirus 1, Equine
- Immunoglobulins against feline Calicivirus, strain 255, Equine
- Immunoglobulins against feline Calicivirus, strain 2024, Equine

Product identification

Medicine name:
Feliserin Plus Injektionslösung für Katzen und Hunde

Active substance:
HORSE PROTEINS
Immunoglobulins against feline panleukopenia virus, Equine
Immunoglobulins against Felid herpesvirus 1, Equine
Immunoglobulins against feline Calicivirus, strain 255, Equine
Immunoglobulins against feline Calicivirus, strain 2024, Equine

Target species:
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
HORSE PROTEINS
50.00 milligram(s) / 1.00 millilitre(s)

Immunoglobulins against feline panleukopenia virus, Equine
10000.00 50% neutralisation dose / 1.00 millilitre(s)

Immunoglobulins against Felid herpesvirus 1, Equine
100.00 50% neutralisation dose / 1.00 millilitre(s)

Immunoglobulins against feline Calicivirus, strain 255, Equine
10000.00 50% neutralisation dose / 1.00 millilitre(s)

Immunoglobulins against feline Calicivirus, strain 2024, Equine
10000.00 50% neutralisation dose / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI06AM01

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

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
Available only in German

Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Tiergesundheit GmbH

Marketing authorisation date:
30/01/2004

Manufacturing sites for batch release:
Ceva Tiergesundheit (Riems) GmbH
IDT Biologika GmbH

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
35a/92

Date of authorisation status change:
14/04/2009

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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