Vetemex 10 mg/ml solution for injection for dogs and cats

Authorised

- Maropitant

Product identification

Medicine name:
Vetemex 10 mg/ml solution for injection for dogs and cats
Vetemex, 10mg/ml, Injekční roztok

Active substance:
Maropitant

Target species:
Dog
Cat

Route of administration:
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Maropitant
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Dog**
- **Cat**

Subcutaneous use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA04AD90

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Czechia

Available in:

Czechia

Package description:

(ID4) 50 millilitre(s): unspecified outer container with 1 Vial (Glass) with 50 millilitre(s)
(ID3) 25 millilitre(s): unspecified outer container with 1 Vial (Glass) with 25 millilitre(s)
(ID2) 20 millilitre(s): unspecified outer container with 1 Vial (Glass) with 20 millilitre(s)
(ID1) 10 millilitre(s): unspecified outer container with 1 Vial (Glass) with 10 millilitre(s)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:
25/02/2019

Manufacturing sites for batch release:
Cp-Pharma Handelsgesellschaft mbH

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicines

Authorisation number:
96/010/19-C

Date of authorisation status change:
25/02/2019

Reference member state:
Germany

Procedure number:
DE/V/0304/001

Concerned member states:
Austria Belgium Croatia Czechia Denmark Estonia Finland France Greece Hungary Ireland Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

Summary of Product Characteristics

English (PDF)
Published on: 10/08/2022
Download

Package Leaflet

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Labelling

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