Vominil 10 mg/ml Injektionslösung für Hunde und Katzen

Authorised

- Maropitant citrate monohydrate

Product identification

Medicine name:
Vominil 10 mg/ml Injektionslösung für Hunde und Katzen
Vominil 10 mg/ml ενέσιμο διάλυμα για σκύλους και γάτες

Active substance:
Maropitant citrate monohydrate

Target species:
Dog
Cat

Route of administration:
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Maropitant citrate monohydrate
14.48 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:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Greece

Package description:
Amber glass vial type I (Ph. Eur.) with 10 ml solution for injection, closed with a chlorobutyl rubber stopper, type I (Ph. Eur) and aluminium pull or flip off cap in a cardboard box.
Amber glass vial type I (Ph. Eur.) with 25 ml solution for injection, closed with a chlorobutyl rubber stopper, type I (Ph. Eur) and aluminium pull or flip off cap in a cardboard box.
Amber glass vial type I (Ph. Eur.) with 50 ml solution for injection, closed with a chlorobutyl rubber stopper, type I (Ph. Eur) and aluminium pull or flip off cap in a cardboard box.
5 x Amber glass vials type I (Ph. Eur.) with 10 ml solution for injection, closed with a chlorobutyl rubber stopper, type I (Ph. Eur) and aluminium pull or flip off cap in a cardboard box.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Vetviva Richter GmbH

Marketing authorisation date:
28/09/2023

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
National Organization For Medicines

Authorisation number:
99122/29-09-2023/K-0255401

Date of authorisation status change:
28/09/2023

Reference member state:
Austria

Procedure number:
AT/V/0030/001

Concerned member states:
Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Netherlands Norway 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

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