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Vominil 10 mg/ml Solution for Injection for Dogs and Cats

Authorised

• Maropitant citrate monohydrate

Product identification

Medicine name:

Vominil 10 mg/ml Injektionslösung für Hunde und Katzen Vominil 10 mg/ml Solution for Injection for Dogs and Cats

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:

OA04AD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Available in:

United Kingdom (Northern Ireland)

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:

29/09/2023

Manufacturing sites for batch release:

Vetviva Richter GmbH

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 57446/3004

Date of authorisation status change:

23/04/2024

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