

Emevet 24 mg - Chewable tablet

Authorised

- Maropitant

Product identification

Medicine name:

Emevet 24 mg - Chewable tablet

Active substance:

Maropitant

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Maropitant

24.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA04AD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

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)

Available in:

Belgium , Czechia , Denmark , Estonia , Finland , Germany , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Norway , Portugal , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:blister (aluminium), Package_size:100 tablets
Packaging:blister (aluminium), Package_size:10 tablets
Packaging:blister (aluminium), Package_size:30 tablets
Packaging:blister (aluminium), Package_size:50 tablets
Packaging:blister (aluminium), Package_size:4 tablets
Packaging:blister (aluminium), Package_size:12 tablets
Packaging:blister (aluminium), Package_size:40 tablets

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:

CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:

2/06/2025

Manufacturing sites for batch release:

CP-Pharma Handelsgesellschaft mbH

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

2/06/2025

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

Documents

Combined File of all Documents

English (PDF)

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