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Emevet 60 mg - Chewable tablet

Authorised

- Maropitant

Product identification

Medicine name:
Emevet 60 mg - Chewable tablet

Active substance:
Maropitant

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Maropitant
60.00 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 , Croatia , Czechia , Denmark , Estonia , Finland , Germany , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Norway , Poland , Portugal , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:
Packaging:blister (aluminium), Package_size:30 tablets
Packaging:blister (aluminium), Package_size:50 tablets
Packaging:blister (aluminium), Package_size:10 tablets
Packaging:blister (aluminium), Package_size:100 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)
Published on: 14/03/2026
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