

# Cimalgex 30 mg - Chewable tablet

Authorised

- Cimicoxib

## Product identification

**Medicine name:**

Cimalgex 30 mg - Chewable tablet

---

**Active substance:**

Cimicoxib

---

**Target species:**

Dog

---

**Route of administration:**

Oral use

---

## Product details

**Active substance and strength:**

Cimicoxib

30.00 milligram(s) / 1.00 Tablet

---

**Pharmaceutical form:**

Chewable tablet

---

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QM01AH93

---

**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:**

Austria , Belgium , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

---

**Package description:**

Packaging: Bottle (HDPE) with child-resistant closure (PP), Package\_size: 45 tablets

Packaging: Blister (Alu/Alu), Package\_size: 144 tablets

Packaging: Blister (Alu/Alu), Package\_size: 32 tablets

Packaging: Blister (Alu/Alu), Package\_size: 8 tablets

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Vetoquinol SA

---

**Marketing authorisation date:**

18/02/2011

---

**Manufacturing sites for batch release:**

Vetoquinol S.A.

---

**Responsible authority:**

European Commission

---

**Authorisation number:**

This information is not available for this product.

---

**Date of authorisation status change:**

18/02/2011

---

To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

Published on: 6/11/2024

[Download](#)

ema-puar-cimalgex-v-162-par-en.pdf