

File downloaded on 2026-04-10

**Source URL:** <https://medicines.health.europa.eu/veterinary/en/600000000609>

# Previcox 57 mg - Chewable tablet

Authorised

- Firocoxib

## Product identification

**Medicine name:**

Previcox 57 mg - Chewable tablet

---

**Active substance:**

Firocoxib

---

**Target species:**

Dog

---

**Route of administration:**

Oral use

---

## Product details

**Active substance and strength:**

Firocoxib

56.88 milligram(s) / 1.00 Tablet

---

**Pharmaceutical form:**

Chewable tablet

---

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

QM01AH90

---

**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 , Estonia , France , Latvia , Lithuania , Luxembourg , Poland , Slovakia , Spain

---

**Package description:**

Packaging:Bottle (PET), Package\_size:60 tablets

Packaging:Blister (Alu/PVC), Package\_size:180 tablets

Packaging:Blister (Alu/PVC), Package\_size:30 tablets

Packaging:Blister (Alu/PVC), Package\_size:10 tablets

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Complete application (stand-alone) - Council Directive 81/851/EEC

---

**Marketing authorisation holder:**

Boehringer Ingelheim Vetmedica GmbH

---

**Marketing authorisation date:**

13/09/2004

---

**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France SCS

---

**Responsible authority:**

European Commission

---

**Authorisation number:**

This information is not available for this product.

---

**Date of authorisation status change:**

19/06/2014

---

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: 29/02/2024

[Download](#)

ema-puar-previcox-v-082-var-ii-0028-en.pdf

ema-puar-previcox-v-082-par-en.pdf