# Rogiola 6 mg chewable tablets for cats

Authorised

- Robenacoxib

## Product identification

**Medicine name:**
Rogiola 6 mg chewable tablets for cats

**Active substance:**
Robenacoxib

**Target species:**
Cat

**Route of administration:**
Oral use

## Product details

**Active substance and strength:**
Robenacoxib
6.00 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**
Chewable tablet

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QM01AH91

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
France

**Package description:**
OPA/Al/PVC/Aluminium blisters containing 6 tablets: 6 x 1 chewable tablet in perforated unit dose blisters, in a cardboard box.
OPA/Al/PVC/Aluminium blisters containing 10 tablets: 10 x 1 chewable tablet in perforated unit dose blisters, in a cardboard box.
OPA/Al/PVC/Aluminium blisters containing 10 tablets: 30 x 1 chewable tablet in perforated unit dose blisters, in a cardboard box.
OPA/Al/PVC/Aluminium blisters containing 10 tablets: 60 x 1 chewable tablet in perforated unit dose blisters, 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:**
KRKA tovarna zdravil d.d. Novo mesto

**Marketing authorisation date:**
28/01/2026

**Manufacturing sites for batch release:**
TAD Pharma GmbH
Krka-Farma d.o.o.

KRKA tovarna zdravil d.d. Novo mesto

**Responsible authority:**
French Agency For Food, Environmental And Occupational Health & Safety

**Authorisation number:**
FR/V/5283679 6/2025

**Date of authorisation status change:**
28/01/2026

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0785/005

**Concerned member states:**
Austria Belgium Cyprus Denmark Finland France Germany Greece Italy Netherlands Norway Portugal Spain Sweden

**Generic of:**
600000004401

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

## Documents

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

English (PDF)
Published on: 26/04/2026
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Combined File of all Documents

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