# Rogiola 10 mg chewable tablets for dogs

Authorised

Robenacoxib

## Product identification

#### **Medicine name:**

Rogiola 10 mg chewable tablets for dogs

Rogiola 10 mg comprimés à croquer pour chiens

Rogiola 10 mg kauwtabletten voor honden

Rogiola 10 mg Kautabletten für Hunde

#### **Active substance:**

Robenacoxib

## **Target species:**

Dog

#### Route of administration:

Oral use

## **Product details**

## **Active substance and strength:**

Robenacoxib

10.00 milligram(s) / 1.00 Tablet

#### **Pharmaceutical form:**

Chewable tablet

## Withdrawal period by route of administration: Oral use:

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Dog

## Anatomical therapeutic chemical veterinary (ATCvet) codes:

OM01AH91

#### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### **Authorised in:**

Belgium

#### Package description:

OPA/Al/PVC/Aluminium perforated blister containing 10 tablets:  $60 \times 1$  chewable tablet in perforated unit dose blisters, in a cardboard box.

OPA/Al/PVC/Aluminium perforated blister containing 10 tablets:  $30 \times 1$  chewable tablet in perforated unit dose blisters, in a cardboard box.

OPA/Al/PVC/Aluminium perforated blister containing 10 tablets:  $10 \times 1$  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:

12/09/2023

## Manufacturing sites for batch release:

TAD Pharma GmbH

Krka-Farma d.o.o.

KRKA tovarna zdravil d.d. Novo mesto

### **Responsible authority:**

Federal Agency For Medicines And Health Products

#### **Authorisation number:**

BE-V661747

#### Date of authorisation status change:

12/09/2023

#### **Reference member state:**

Ireland

#### **Procedure number:**

IE/V/0785/002

#### **Concerned member states:**

Austria Belgium Cyprus Denmark Finland France Germany Greece Italy Netherlands Norway Portugal Spain Sweden

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

## **Documents**

Summary of Product Characteristics

English (PDF)

Published on: 3/05/2024

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Package Leaflet

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Labelling

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