# Robexera 40 mg chewable tablets for dogs

• Robenacoxib

# **Product identification**

## Medicine name:

Robexera 40 mg chewable tablets for dogs Robexera 40 mg таблетки за дъвчене за кучета

## Active substance:

Robenacoxib

### **Target species:**

Dog

# Route of administration:

Oral use

# **Product details**

## Active substance and strength:

Robenacoxib 40.00 milligram(s) / 1.00 Tablet

## **Pharmaceutical form:**

Chewable tablet

## Withdrawal period by route of administration:

#### Oral use:

Dog

## Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AH91

## Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

### Authorisation status:

Valid

## Authorised in:

Bulgaria

## Package description:

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

OPA/AI/PVC/Aluminium perforated blister containing 10 tablets: 30 x 1 chewable tablet in perforated unit dose blisters, in a cardboard box.

OPA/Al/PVC/Aluminium perforated blister containing 10 tablets: 10 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:

20/06/2023

## Manufacturing sites for batch release:

Krka-Farma d.o.o. TAD Pharma GmbH KRKA tovarna zdravil d.d. Novo mesto

## Responsible authority:

Bulgarian Food Safety Authority

Authorisation number: 0022-3193

**Date of authorisation status change:** 20/06/2023

20/00/2025

Reference member state:

Ireland

## Procedure number:

IE/V/0775/004

### **Concerned member states:**

Belgium Bulgaria Croatia Czechia Denmark Estonia Finland France Germany Hungary Italy Latvia Lithuania Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

## Documents

Summary of Product Characteristics

English (PDF)

Package Leaflet

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

Labelling

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

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