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# Robexera 10 mg chewable tablets for dogs

Authorised

Robenacoxib

# Product identification

#### **Medicine name:**

Robexera 10 mg chewable tablets for dogs Robexera, 10 mg, kramtomosios tabletės šunims

#### **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

## Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AH91

## Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### **Authorised in:**

Lithuania

#### **Available in:**

Lithuania

## 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:

1/08/2023

# Manufacturing sites for batch release:

Krka-Farma d.o.o.

TAD Pharma GmbH KRKA tovarna zdravil d.d. Novo mesto

## **Responsible authority:**

State Food And Veterinary Service

#### **Authorisation number:**

LT/2/23/2763/001-003

## Date of authorisation status change:

1/08/2023

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

Ireland

## **Procedure number:**

IE/V/0775/002

#### **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)

Published on: 3/05/2024

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