

# Robexera 20 mg chewable tablets for dogs

Authorised

- Robenacoxib

## Product identification

**Medicine name:**

Robexera 20 mg chewable tablets for dogs  
Robexera 20 mg Tabletka do rozgryzania i żucia

**Active substance:**

Robenacoxib

**Target species:**

Dog

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Robenacoxib  
20.00 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**

Chewable tablet

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

QM01AH91

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Poland

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**Available in:**

Poland

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**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/Al/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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 18 of Regulation (EU) 2019/6)

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**Marketing authorisation holder:**

KRKA tovarna zdravil d.d. Novo mesto

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**Marketing authorisation date:**

25/07/2023

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**Manufacturing sites for batch release:**

Krka-Farma d.o.o.

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

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

3261

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**Date of authorisation status change:**

25/07/2023

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0775/003

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**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)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

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.

## Package Leaflet

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