# Robexera 40 mg chewable tablets for dogs

Authorised

Robenacoxib

# Product identification

#### **Medicine name:**

Robexera 40 mg chewable tablets for dogs Robexera 40 mg comprimidos mastigáveis para cães

### **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:

Portugal

## 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:

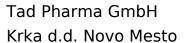
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# Marketing authorisation date:

14/07/2023

# Manufacturing sites for batch release:

Krka-Farma d.o.o.



## **Responsible authority:**

Directorate General For Food And Veterinary

#### **Authorisation number:**

1586/04/23DFVPT

## Date of authorisation status change:

14/07/2023

#### Reference member state:

Ireland

#### **Procedure number:**

IE/V/0775/004

## **Concerned member states:**

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

To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>

## **Documents**

Combined File of all Documents

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| Summary of Product Characteristics |  |
|------------------------------------|--|
| Summary of Product Characteristics |  |

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