

Robexera 20 mg/ml solution for injection for cats and dogs

Authorised

- Robenacoxib

Product identification

Medicine name:

Robexera 20 mg/ml solution for injection for cats and dogs

Active substance:

Robenacoxib

Target species:

Dog

Cat

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Robenacoxib

20.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AH91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Estonia

Package description:

A cardboard box containing one amber type I glass vial of 20 ml, closed with a type I bromobutyl rubber stopper and an aluminium seal with plastic tear-off tab.

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:

27/08/2025

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

TAD Pharma GmbH

Responsible authority:

State Agency Of Medicines

Authorisation number:

1211425

Date of authorisation status change:

27/08/2025

Reference member state:

Ireland

Procedure number:

IE/V/0627/001

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)

Generic of:

600000004139

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 24/08/2025

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Combined File of all Documents

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