Robexera 20 mg chewable tablets for dogs

Authorised

- Robenacoxib

Product identification

Medicine name:
Robexera 20 mg chewable tablets for dogs
Robexera vet 20 mg tuggtablett för hund

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

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:

Sweden

Package description:

Available only in Swedish

Available only in Swedish

Available only in Swedish

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:

8/09/2023

Manufacturing sites for batch release:

Krka-Farma d.o.o.

Tad Pharma GmbH

Krka d.d. Novo Mesto

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
63436

Date of authorisation status change:
8/09/2023

Reference member state:
Ireland

Procedure number:
IE/V/0775/003

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 www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)
Published on: 3/05/2024
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Package Leaflet

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Labelling

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