

Rycarfa 20 mg tablets for dogs

Authorised

- Carprofen

Product identification

Medicine name:

Rycarfa 20 mg tablets for dogs

Rycarfa 20 mg Tablet

Rycarfa 20 mg Comprimé

Rycarfa 20 mg Tablette

Active substance:

Carprofen

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Carprofen

20.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Withdrawal period by route of administration:

Oral use:

-

Dog

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AE91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Package description:

Blister (OPA/Al/PVC-Al): 20 tablets (10 tablets/blister) in a box.

Blister (OPA/Al/PVC-Al): 50 tablets (10 tablets/blister) in a box.

Blister (OPA/Al/PVC-Al): 100 tablets (10 tablets/blister) in a box.

Blister (OPA/Al/PVC-Al): 500 tablets (10 tablets/blister) in a box.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:

27/03/2014

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

TAD Pharma GmbH
Krka-Farma d.o.o.

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V456773

Date of authorisation status change:

27/03/2014

Reference member state:

Ireland

Procedure number:

IE/V/0316/001

Concerned member states:

Austria Belgium Denmark Finland France Germany Greece Italy
Netherlands Norway Portugal Spain 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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