Rycarfa 50 mg tablets for dogs

Authorised

- Carprofen

Product identification

Medicine name:
Rycarfa 50 mg tablets for dogs

Active substance:
Carprofen

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Carprofen
50.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AE91

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Netherlands

Available in:
Netherlands

Package description:
Blister (OPA/Al/PVC-Al): 500 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): 50 tablets (10 tablets/blister) in a box
Blister (OPA/Al/PVC-Al): 20 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:
17/12/2013

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto
Krka-Farma d.o.o.
TAD Pharma GmbH

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 112845

Date of authorisation status change:
26/01/2022

Reference member state:
Ireland

Procedure number:
IE/V/0316/002

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

Combined File of all Documents

English (PDF)
Published on: 24/08/2025
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Summary of Product Characteristics