Karprovet 50 mg tablets for dogs

Authorised

- Carprofen

Product identification

Medicine name:
Karprovet 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:
France

Available in:
France

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:
7/05/2010

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

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/5807577 7/2010

Date of authorisation status change:
16/07/2015

Reference member state:
Ireland

Procedure number:
IE/V/0239/003

Concerned member states:
Belgium Czechia Estonia France Germany Hungary Italy Latvia Lithuania Poland Portugal Romania Slovakia Slovenia 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: 24/08/2025
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Combined File of all Documents

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.