File downloaded on 2026-11-26
Source URL: https://medicines.health.europa.eu/veterinary/en/600000047011

Rycarfa 50 mg/ml solution for injection for dogs and cats

Authorised

- Carprofen

Product identification

Medicine name:
Rycarfa 50 mg/ml solution for injection for dogs and cats

Active substance:
Carprofen

Target species:
Dog
Cat

Route of administration:
Intravenous use
Subcutaneous use

Product details

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

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AE91

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

Authorisation status:
Valid

Authorised in:
France

Package description:
Type I glass vial (amber glass): 1 vial of 20 ml solution for injection with bromobutyl rubber stopper and aluminium seal, 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:
3/02/2014

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

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

Authorisation number:

FR/V/3231402 9/2014

Date of authorisation status change:
22/12/2021

Reference member state:
Ireland

Procedure number:
IE/V/0317/001

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: 25/09/2024
Updated on: 14/03/2026
Download

Package Leaflet and Labelling

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