File downloaded on 2026-10-30
Source URL: https://medicines.health.europa.eu/veterinary/en/600000047045

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:
Portugal

Available in:
Portugal

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:
30/12/2013

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

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:
750/01/13DFVPT

Date of authorisation status change:
29/10/2024

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

Combined File of all Documents

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