

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

Rycarfa 50 mg/ml oplossing voor injectie voor honden en katten

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:

Netherlands

Available in:

Netherlands

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:

27/01/2014

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

TAD Pharma GmbH

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 112830

Date of authorisation status change:

26/01/2022

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

Combined File of all Documents

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

Summary of Product Characteristics