

Carporal VET. 40 mg tablets for dogs

Authorised

- Carprofen

Product identification

Medicine name:

Carporal VET. 40 mg tablets for dogs

Canicaral vet 40 mg Tabletka

Active substance:

Carprofen

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Carprofen

40.00 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:

Poland

Package description:

Cardboard box of 50 blisters (Aluminium - PA/ALU/PVC) of 10 tablets

Cardboard box of 5 blisters (Aluminium - PA/ALU/PVC) of 10 tablets

Cardboard box of 4 blisters (Aluminium - PA/ALU/PVC) of 10 tablets

Cardboard box of 25 blisters (Aluminium - PA/ALU/PVC) of 10 tablets

Cardboard box of 3 blisters (Aluminium - PA/ALU/PVC) of 10 tablets

Cardboard box of 2 blisters (Aluminium - PA/ALU/PVC) of 10 tablets

Cardboard box of 10 blisters (Aluminium - PA/ALU/PVC) of 10 tablets

Cardboard box of 1 blister (Aluminium - PA/ALU/PVC) of 10 tablets

Cardboard box of 6 blisters (Aluminium - PA/ALU/PVC) of 10 tablets

Cardboard box of 8 blisters (Aluminium - PA/ALU/PVC) of 10 tablets

Cardboard box of 7 blisters (Aluminium - PA/ALU/PVC) of 10 tablets

Cardboard box of 9 blisters (Aluminium - PA/ALU/PVC) of 10 tablets

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Le Vet. Beheer B.V.

Marketing authorisation date:

13/01/2020

Manufacturing sites for batch release:

Artesan Pharma GmbH & Co. KG
Lelypharma B.V.
Genera d.d.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

2932

Date of authorisation status change:

13/01/2020

Reference member state:

Netherlands

Procedure number:

NL/V/0191/001

Concerned member states:

Austria Belgium Croatia Cyprus Czechia Denmark Estonia Finland France
Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Norway
Poland Portugal Romania Slovakia Slovenia Spain Sweden
United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Package Leaflet

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

Summary of Product Characteristics

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

Labelling

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

Combined File of all Documents