Carporal VET. 40 mg tablets for dogs

Authorised

- Carprofen

Product identification

Medicine name:
Carporal VET. 40 mg tablets for dogs

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:
Finland

Available in:
Finland

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:
18/11/2015

Manufacturing sites for batch release:
Artesan Pharma GmbH & Co. KG
Lelypharma B.V.
Genera d.d.

Responsible authority:
Finnish Medicines Agency

Authorisation number:
32437

Date of authorisation status change:
18/11/2015

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

Summary of Product Characteristics

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

Package Leaflet

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

Combined File of all Documents