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Carporal VET. 160 mg tablets for dogs

Authorised

- Carprofen

Product identification

Medicine name:
Carporal VET. 160 mg tablets for dogs

Active substance:
Carprofen

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Carprofen
160.00 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:
Czechia

Package description:
Cardboard box of 25 blisters (Aluminium - PA/ALU/PVC blister) of 10 tablets
Cardboard box of 10 blisters (Aluminium - PA/ALU/PVC blister) of 10 tablets
Cardboard box of 2 blisters (Aluminium - PA/ALU/PVC blister) of 10 tablets
Cardboard box of 1 blister (Aluminium - PA/ALU/PVC blister) of 10 tablets
Cardboard box of 50 blisters (Aluminium - PA/ALU/PVC blister) of 10 tablets
Cardboard box of 6 blisters (Aluminium - PA/ALU/PVC blister) of 10 tablets
Cardboard box of 7 blisters (Aluminium - PA/ALU/PVC blister) of 10 tablets
Cardboard box of 3 blisters (Aluminium - PA/ALU/PVC blister) of 10 tablets
Cardboard box of 4 blisters (Aluminium - PA/ALU/PVC blister) of 10 tablets
Cardboard box of 5 blisters (Aluminium - PA/ALU/PVC blister) of 10 tablets
Cardboard box of 8 blisters (Aluminium - PA/ALU/PVC blister) of 10 tablets
Cardboard box of 9 blisters (Aluminium - PA/ALU/PVC blister) 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:

8/03/2016

Manufacturing sites for batch release:
Genera d.d.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/017/16-C

Date of authorisation status change:
26/08/2020

Reference member state:
Netherlands

Procedure number:
NL/V/0191/002

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

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Labelling

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.