

# Carporal VET. 160 mg tablets for dogs

Authorised

- Carprofen

## Product identification

**Medicine name:**

Carporal VET. 160 mg tablets for dogs

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**Active substance:**

Carprofen

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Carprofen

160.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QM01AE91

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Lithuania

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**Package description:**

Cardboard box of 9 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 5 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 3 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 6 blisters (Aluminium - PA/ALU/PVC blister) of 10 tablets  
Cardboard box of 50 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 2 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 25 blisters (Aluminium - PA/ALU/PVC blister) of 10 tablets

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Le Vet. Beheer B.V.

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**Marketing authorisation date:**

27/07/2015

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**Manufacturing sites for batch release:**

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

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**Responsible authority:**

State Food And Veterinary Service

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**Authorisation number:**

LT/2/15/2297/001-012

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**Date of authorisation status change:**

13/10/2020

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0191/002

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**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)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

RV2297.pdf