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# CARPRODYL QUADRI 120 MG CHEWABLE TABLETS FOR DOGS

Authorised

Carprofen

# Product identification

#### **Medicine name:**

CARPRODYL QUADRI 120 MG CHEWABLE TABLETS FOR DOGS Carprodyl Quadri, 120 mg närimistabletid koertele

### **Active substance:**

Carprofen

# **Target species:**

Dog

#### Route of administration:

Oral use

# **Product details**

## **Active substance and strength:**

Carprofen

120.00 milligram(s) / 1.00 Tablet

#### **Pharmaceutical form:**

Chewable tablet

Authorised in: Estonia  Package description: Cardboard box with 2 blisters of 6 tablets Cardboard box with 80 blisters of 6 tablets Cardboard box with 40 blisters of 6 tablets Cardboard box with 20 blisters of 6 tablets	
Authorisation status: Valid  Authorised in: Estonia  Package description: Cardboard box with 2 blisters of 6 tablets Cardboard box with 80 blisters of 6 tablets Cardboard box with 40 blisters of 6 tablets Cardboard box with 20 blisters of 6 tablets	
Package description: Cardboard box with 2 blisters of 6 tablets Cardboard box with 80 blisters of 6 tablets Cardboard box with 40 blisters of 6 tablets Cardboard box with 20 blisters of 6 tablets	
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Package description: Cardboard box with 2 blisters of 6 tablets Cardboard box with 80 blisters of 6 tablets Cardboard box with 40 blisters of 6 tablets Cardboard box with 20 blisters of 6 tablets Additional information	
Cardboard box with 2 blisters of 6 tablets Cardboard box with 80 blisters of 6 tablets Cardboard box with 40 blisters of 6 tablets Cardboard box with 20 blisters of 6 tablets	
Cardboard box with 80 blisters of 6 tablets Cardboard box with 40 blisters of 6 tablets Cardboard box with 20 blisters of 6 tablets	
Cardboard box with 40 blisters of 6 tablets	
Cardboard box with 20 blisters of 6 tablets	
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:	
Ceva Sante Animale	
Marketing authorisation date: 27/02/2017	
Manufacturing sites for batch release: CEVA Santé Animale	

**Authorisation number:** 

# Date of authorisation status change:

27/02/2017

### Reference member state:

France

## **Procedure number:**

FR/V/0172/002

## **Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands 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

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