

# Prednicortone 5 mg tablets for dogs and cats

Authorised

- Prednisolone

## Product identification

**Medicine name:**

Prednicortone 5 mg tablets for dogs and cats

Prednicortone, 5 mg, tableta, za pse i mačke

**Active substance:**

Prednisolone

**Target species:**

Dog

Cat

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Prednisolone

5.00 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**

Tablet

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

QH02AB06

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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:**

Croatia

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**Package description:**

Cardboard box of 1 Aluminium - PVC/PE/PVDC blister of 10 tablets.  
Cardboard box of 5 Aluminium - PVC/PE/PVDC blisters of 10 tablets.  
Cardboard box of 4 Aluminium - PVC/PE/PVDC blisters of 10 tablets.  
Cardboard box of 3 Aluminium - PVC/PE/PVDC blisters of 10 tablets.  
Cardboard box of 15 Aluminium - PVC/PE/PVDC blisters of 10 tablets.  
Cardboard box of 25 Aluminium - PVC/PE/PVDC blisters of 10 tablets.  
Cardboard box of 2 Aluminium - PVC/PE/PVDC blisters of 10 tablets.  
Cardboard box of 10 Aluminium - PVC/PE/PVDC blisters of 10 tablets.  
Cardboard box of 8 Aluminium - PVC/PE/PVDC blisters of 10 tablets.  
Cardboard box of 7 Aluminium - PVC/PE/PVDC blisters of 10 tablets.  
Cardboard box of 6 Aluminium - PVC/PE/PVDC blisters of 10 tablets.  
Cardboard box of 50 Aluminium - PVC/PE/PVDC blisters of 10 tablets.  
Cardboard box of 9 Aluminium - PVC/PE/PVDC blisters of 10 tablets.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Dechra Regulatory B.V.

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**Marketing authorisation date:**

13/10/2016

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**Manufacturing sites for batch release:**

Lelypharma B.V.

Genera d.d.

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**Responsible authority:**

Ministry Of Agriculture Veterinary And Food Safety Directorate

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**Authorisation number:**

UP/I-322-05/20-01/338

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**Date of authorisation status change:**

26/03/2025

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0190/001

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**Concerned member states:**

Austria Belgium Croatia Cyprus Czechia Denmark Estonia Finland France

Germany 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

Summary of Product Characteristics

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Combined File of all Documents