

# Dexacortone 0,5 mg tablets for dogs and cats

Authorised

- Dexamethasone

## Product identification

**Medicine name:**

Dexacortone 0,5 mg tablets for dogs and cats

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**Active substance:**

Dexamethasone

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**Target species:**

Dog

Cat

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Dexamethasone

0.50 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Chewable tablet

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

QH02AB02

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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:**

Finland

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**Available in:**

Finland

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**Package description:**

Aluminium - PVC-PE-PVDC blister. Cardboard box of 2 blisters of 10 tablets.

Aluminium - PVC-PE-PVDC blister. Cardboard box of 6 blisters of 10 tablets.

Aluminium - PVC-PE-PVDC blister. Cardboard box of 5 blisters of 10 tablets.

Aluminium - PVC-PE-PVDC blister. Cardboard box of 1 blister of 10 tablets.

Aluminium - PVC-PE-PVDC blister. Cardboard box of 9 blisters of 10 tablets.

Aluminium - PVC-PE-PVDC blister. Cardboard box of 4 blisters of 10 tablets.

Aluminium - PVC-PE-PVDC blister. Cardboard box of 10 blisters of 10 tablets.

Aluminium - PVC-PE-PVDC blister. Cardboard box of 3 blisters of 10 tablets.

Aluminium - PVC-PE-PVDC blister. Cardboard box of 8 blisters of 10 tablets.

Aluminium - PVC-PE-PVDC blister. Cardboard box of 7 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:**

Le Vet. Beheer B.V.

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**Marketing authorisation date:**

11/12/2022

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**Manufacturing sites for batch release:**

Lelypharma B.V.

Genera d.d.

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**Responsible authority:**

Finnish Medicines Agency

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**Authorisation number:**

34793

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**Date of authorisation status change:**

11/12/2022

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0219/001

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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

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