

# 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:**

Belgium

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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:**

20/02/2018

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**Manufacturing sites for batch release:**

Lelypharma B.V.

Genera d.d.

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**Responsible authority:**

Federal Agency For Medicines And Health Products

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**Authorisation number:**

BE-V525840

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**Date of authorisation status change:**

20/02/2018

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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.

## Labelling

This document does not exist in this language (English). You can find it in another language below.

## Combined File of all Documents