

# DERMIPRED 10 MG TABLETS FOR DOGS

Authorised

- Prednisolone

## Product identification

**Medicine name:**

DERMIPRED 10 MG TABLETS FOR DOGS

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**Active substance:**

Prednisolone

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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:**

Prednisolone

10.00 milligram(s) / 1.00 Tablet

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**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:**

Finland

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**Package description:**

Cardboard box with 16 tablets (Aluminium / Polyvinylidene chloride - Thermo elast - Polyvinyl chloride blister)

Cardboard box with 96 tablets (Aluminium / Polyvinyl chloride - Aluminium - Polyamide blister)

Cardboard box with 16 tablets (Aluminium / Polyvinyl chloride - Aluminium - Polyamide blister)

Cardboard box with 96 tablets (Aluminium / Polyvinylidene chloride - Thermo elast - Polyvinyl chloride blister)

Cardboard box with 15 tablets (Aluminium / Polyvinyl chloride - Aluminium - Polyamide blister)

Cardboard box with 90 tablets (Aluminium / Polyvinyl chloride - Aluminium - Polyamide blister)

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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:**

Ceva Sante Animale

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**Marketing authorisation date:**

21/12/2016

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**Manufacturing sites for batch release:**

Ceva Sante Animale

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**Responsible authority:**

Finnish Medicines Agency

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**Authorisation number:**

33506

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**Date of authorisation status change:**

21/12/2016

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**Reference member state:**

France

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**Procedure number:**

FR/V/0301/002

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**Concerned member states:**

Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland  
Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg  
Norway Poland Portugal Romania Slovakia 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

eu-puar-frv0301002-mr-rpe425-en.pdf