

DERMIPRED 5 MG TABLETS FOR DOGS

Authorised

- Prednisolone

Product identification

Medicine name:

DERMIPRED 5 MG TABLETS FOR DOGS

Active substance:

Prednisolone

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Prednisolone

5.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Sweden

Available in:

Sweden

Package description:

Available only in [Swedish](#)

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Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Ceva Sante Animale

Marketing authorisation date:

5/10/2016

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

Swedish Medical Products Agency

Authorisation number:

53215

Date of authorisation status change:

5/10/2016

Reference member state:

France

Procedure number:

FR/V/0301/001

Concerned member states:

Austria Belgium Bulgaria Cyprus Czechia Denmark Finland Germany
Greece Hungary Ireland Italy Netherlands Norway Poland Portugal Romania
Slovakia Spain Sweden United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Package Leaflet

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

Summary of Product Characteristics

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

eu-puar-frv0301001-mr-rpe241-en.pdf