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DERMIPRED 10 MG TABLETS FOR DOGS

Authorised

- Prednisolone

Product identification

Medicine name:
DERMIPRED 10 MG TABLETS FOR DOGS

Active substance:
Prednisolone

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Prednisolone
10.00 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:
Poland

Available in:
Poland

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)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Ceva Animal Health Polska Sp. z o.o.

Marketing authorisation date:
14/04/2017

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2650

Date of authorisation status change:
14/04/2017

Reference member state:
France

Procedure number:
FR/V/0301/002

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)

To consult adverse reactions on veterinary medicinal products please go to 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.

Labelling

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