

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:

Hungary

Available in:

Hungary

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-Phylaxia Zrt.

Marketing authorisation date:

18/09/2018

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

3983/X/18 NÉBIH ÁTI (16 tableta aluminium/polivin

Date of authorisation status change:

18/09/2018

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

Combined File of all Documents

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