

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:

Denmark

Available in:

Denmark

Package description:

Cardboard box with 120 tablets (Aluminium / Polyvinylidene chloride - Thermo elast - Polyvinyl chloride blister)

Cardboard box with 120 tablets (Aluminium / Polyvinyl chloride - Aluminium - Polyamide blister)

Cardboard box with 24 tablets (Aluminium / Polyvinylidene chloride - Thermo elast - Polyvinyl chloride blister)

Cardboard box with 20 tablets (Aluminium / Polyvinyl chloride - Aluminium - Polyamide blister)

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:

17/08/2016

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

Danish Medicines Agency

Authorisation number:

56309

Date of authorisation status change:

17/08/2016

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

Summary of Product Characteristics

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