

DERMIPRED 10 MG TABLETS FOR DOGS

Authorised

- Prednisolone

Product identification

Medicine name:

DERMIPRED 10 MG TABLETS FOR DOGS
Dermipred 10 mg tableta kutyáknak A.U.V.

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

Withdrawal period by route of administration:

Oral use:

- Dog

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Hungary

Package description:

Cardboard box with 96 tablets

Cardboard box with 16 tablets

Cardboard box with 96 tablets

Cardboard box with 16 tablets

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:

National Food Chain Safety Office

Authorisation number:

3983/X/18 NÉBIH ÁTI (16 tabletta 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 Czechia Denmark Estonia Finland Germany
Hungary Ireland Italy Latvia Lithuania Luxembourg 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

Source URL: <https://medicines.health.europa.eu/veterinary/600000045769>