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DERMIPRED 20 MG TABLETS FOR DOGS

Authorised

- Prednisolone

Product identification

Medicine name:
DERMIPRED 20 MG TABLETS FOR DOGS

Active substance:
Prednisolone

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Prednisolone
20.00 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 20 tablets (Aluminium / Polyvinylidene chloride - Thermo elast - Polyvinyl chloride blister)
Cardboard box with 100 tablets (Aluminium / Polyvinyl chloride - Aluminium - Polyamide blister)
Cardboard box with 20 tablets (Aluminium / Polyvinyl chloride - Aluminium - Polyamide blister)
Cardboard box with 100 tablets (Aluminium / Polyvinylidene chloride - Thermo elast - Polyvinyl chloride 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:
56310

Date of authorisation status change:
17/08/2016

Reference member state:
France

Procedure number:
FR/V/0301/003

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

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