DERMIPRED 20 MG TABLETS FOR DOGS

Authorised

- Prednisolone

Product identification

Medicine name:
DERMIPRED 20 MG TABLETS FOR DOGS
Dermipred 20 mg tabletten voor honden

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

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:
Netherlands

Package description:
Cardboard box with 100 tablets
Cardboard box with 20 tablets
Cardboard box with 100 tablets
Cardboard box with 20 tablets

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 B.V.

Marketing authorisation date:
13/09/2016

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:

Medicines Evaluation Board

Authorisation number:
REG NL 117608

Date of authorisation status change:
28/03/2022

Reference member state:
France

Procedure number:
FR/V/0301/003

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

Documents

Combined File of all Documents

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