Prednizol 5mg Tablets for Dogs and Cats

Authorised

- Prednisolone

Product identification

Medicine name:
Prednizol 5mg Tablets for Dogs and Cats
Prednizol 5 mg tabletten voor honden en katten

Active substance:
Prednisolone

Target species:
Dog
Cat

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:
Netherlands

Package description:
Cardboard box of 1 PVC Aluminium/PVC foil blister of 10 tablets each
White polypropylene container containing 250 tablets, sealed with a white, childresistant, high-density polyethylene closure.
Cardboard box of 2 PVC Aluminium/PVC foil blisters of 10 tablets each
Cardboard box of 3 PVC Aluminium/PVC foil blisters of 10 tablets each
Cardboard box of 4 PVC Aluminium/PVC foil blisters of 10 tablets each
Cardboard box of 5 PVC Aluminium/PVC foil blisters of 10 tablets each
Cardboard box of 6 PVC Aluminium/PVC foil blisters of 10 tablets each
Cardboard box of 7 PVC Aluminium/PVC foil blisters of 10 tablets each
Cardboard box of 8 PVC Aluminium/PVC foil blisters of 10 tablets each
Cardboard box of 9 PVC Aluminium/PVC foil blisters of 10 tablets each
Cardboard box of 10 PVC Aluminium/PVC foil blisters of 10 tablets each
Cardboard box of 12 PVC Aluminium/PVC foil blisters of 10 tablets each
Cardboard box of 15 PVC Aluminium/PVC foil blisters of 10 tablets each
Cardboard box of 25 PVC Aluminium/PVC foil blisters of 10 tablets each
Cardboard box of 50 PVC Aluminium/PVC foil blisters of 10 tablets each

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:
Millpledge Europe

Marketing authorisation date:
9/01/2020

Manufacturing sites for batch release:
Millpledge Europe

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 122989

Date of authorisation status change:
25/01/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0351/001

Concerned member states:
Belgium France Germany 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

English (PDF)

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