Prednizol 5 mg Tablets for Dogs and Cats

Authorised

- Prednisolone

Product identification

Medicine name:
Prednizol 5mg Tablets for Dogs and Cats
Prednizol 5 mg Tablets for Dogs and Cats

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

Withdrawal period by route of administration:

Oral use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

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

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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 Limited

Marketing authorisation date:
10/10/2019

Manufacturing sites for batch release:
Millpledge Europe

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 04409/4008

Date of authorisation status change:
24/08/2022

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
