Dexacortone 0,5 mg tablets for dogs and cats

Authorised

- Dexamethasone

Product identification

Medicine name:
Dexacortone 0,5 mg tablets for dogs and cats
Dexacortone 0,5 mg žuvacie tablety pre psy a mačky

Active substance:
Dexamethasone

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Dexamethasone
0.50 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Chewable tablet

Withdrawal period by route of administration:

Oral use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

Aluminium - PVC-PE-PVDC blister. Cardboard box of 2 blisters of 10 tablets.
Aluminium - PVC-PE-PVDC blister. Cardboard box of 6 blisters of 10 tablets.
Aluminium - PVC-PE-PVDC blister. Cardboard box of 5 blisters of 10 tablets.
Aluminium - PVC-PE-PVDC blister. Cardboard box of 1 blister of 10 tablets.
Aluminium - PVC-PE-PVDC blister. Cardboard box of 9 blisters of 10 tablets.
Aluminium - PVC-PE-PVDC blister. Cardboard box of 4 blisters of 10 tablets.
Aluminium - PVC-PE-PVDC blister. Cardboard box of 10 blisters of 10 tablets.
Aluminium - PVC-PE-PVDC blister. Cardboard box of 3 blisters of 10 tablets.
Aluminium - PVC-PE-PVDC blister. Cardboard box of 8 blisters of 10 tablets.
Aluminium - PVC-PE-PVDC blister. Cardboard box of 7 blisters of 10 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:
Le Vet. Beheer B.V.

Marketing authorisation date:
29/05/2018

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/029/DC/18-S

Date of authorisation status change:
29/05/2018

Reference member state:
Netherlands

Procedure number:
NL/V/0219/001

Concerned member states:
Austria Belgium Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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Documents

Combined File of all Documents

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