

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 vet, 0,5 mg, kramtomosios tabletēs šunims ir katēms

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Lithuania

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:

19/12/2017

Manufacturing sites for batch release:

Lelypharma B.V.
Genera d.d.

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/17/2433/001-010

Date of authorisation status change:

15/12/2025

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)

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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