Lodisure 1 mg tablets for cats

Authorised

- Amlodipine besilate

Product identification

Medicine name:
Lodisure 1 mg tablets for cats

Active substance:
Amlodipine besilate

Target species:
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Amlodipine besilate
1.40 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC08CA01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Poland

Available in:
Poland

Package description:
Blister made of PVC /aluminium / OPA with a push-through PVC-PVDC/aluminium lidding foil. Each blister contains 14 tablets.1 cardboard carton with 56 tablets.
Blister made of PVC /aluminium / OPA with a push-through PVC-PVDC/aluminium lidding foil. Each blister contains 14 tablets.1 cardboard carton with 84 tablets.
Blister made of PVC /aluminium / OPA with a push-through PVC-PVDC/aluminium lidding foil. Each blister contains 14 tablets.1 cardboard carton with 28 tablets.
Blister made of PVC /aluminium / OPA with a push-through PVC-PVDC/aluminium lidding foil. Each blister contains 14 tablets.1 cardboard carton with 168 tablets.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Dechra Regulatory B.V.

Marketing authorisation date:
21/04/2021

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3093

Date of authorisation status change:
21/04/2021

Reference member state:
Netherlands

Procedure number:
NL/V/0339/001

Concerned member states:
Austria Belgium Bulgaria Croatia 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

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.