File downloaded on 2026-11-07
Source URL: https://medicines.health.europa.eu/veterinary/en/600000035439

Amodip 1.25 mg Chewable Tablets for Cats

Authorised

- Amlodipine besilate

Product identification

Medicine name:
Amodip 1.25 mg Chewable Tablets for Cats

Active substance:
Amlodipine besilate

Target species:
Cat

Route of administration:
Oral use

Product details

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

Pharmaceutical form:
Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC08CA01

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

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Available in:
United Kingdom (Northern Ireland)

Package description:
Box containing 3 blister packs of 10 chewable tablets
Box containing 20 blister packs of 10 chewable tablets
Box containing 10 blister packs of 10 chewable 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:
Ceva Sante Animale

Marketing authorisation date:
28/04/2015

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 14966/3021

Date of authorisation status change:
22/06/2024

Reference member state:
France

Procedure number:
FR/V/0413/001

Concerned member states:
Austria Belgium Bulgaria Czechia Denmark Estonia Finland Germany Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands 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

eu-puar-frv0413001-mr-rpe497-en.pdf