

# AMODIP 1.25 MG CHEWABLE TABLETS FOR CATS

Authorised

- Amlodipine besilate

## Product identification

**Medicine name:**

AMODIP 1.25 MG CHEWABLE TABLETS FOR CATS

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**Active substance:**

Amlodipine besilate

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**Target species:**

Cat

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Amlodipine besilate

1.73 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Chewable tablet

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QC08CA01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Denmark

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**Available in:**

Denmark

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**Package description:**

Box containing 10 blister packs of 10 chewable tablets

Box containing 20 blister packs of 10 chewable tablets

Box containing 3 blister packs of 10 chewable tablets

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ceva Sante Animale

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**Marketing authorisation date:**

18/06/2015

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**Manufacturing sites for batch release:**

Ceva Sante Animale

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**Responsible authority:**

Danish Medicines Agency

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**Authorisation number:**

53431

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**Date of authorisation status change:**

18/06/2015

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**Reference member state:**

France

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**Procedure number:**

FR/V/0413/001

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**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)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://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

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