

# Alpramil 16 mg/40 mg film-coated tablets for cats weighing at least 4 kg

Authorised

- Praziquantel
- Milbemyacin oxime

## Product identification

**Medicine name:**

Alpramil 16 mg/40 mg film-coated tablets for cats weighing at least 4 kg

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**Active substance:**

Praziquantel

Milbemyacin oxime

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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:**

Praziquantel

40.00 milligram(s) / 1.00 Tablet

Milbemycin oxime  
16.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Film-coated tablet

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

QP54AB51

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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:**

Croatia

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**Package description:**

Box with 25 PVC / PE / PVDC - Aluminium blisters each containing 2 tablets.

Box with 25 PVC / PE / PVDC - Aluminium blisters each containing 1 tablet.

Box with 10 PVC / PE / PVDC - Aluminium blisters each containing 1 tablet.

Box with 1 PVC / PE / PVDC - Aluminium blister containing 4 tablets.

Box with 25 PVC / PE / PVDC - Aluminium blisters each containing 4 tablets.

Box with 10 PVC / PE / PVDC - Aluminium blisters each containing 4 tablets.

Box with 10 PVC / PE / PVDC - Aluminium blisters each containing 2 tablets.

1 box with 1PVC / PE / PVDC - Aluminium blister containing 2 tablets.

1 box with 1PVC / PE / PVDC - Aluminium blister containing 1 tablet.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Alfasan Nederland B.V.

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**Marketing authorisation date:**

7/04/2022

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**Manufacturing sites for batch release:**

Alfasan Nederland B.V.

Lelypharma B.V.

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**Responsible authority:**

Ministry Of Agriculture Veterinary And Food Safety Directorate

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**Authorisation number:**

UP/I-322-05/22-01/253

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**Date of authorisation status change:**

23/06/2025

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0364/003

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**Concerned member states:**

Austria Belgium Bulgaria 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

Summary of Product Characteristics

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Combined File of all Documents

English (PDF)

Published on: 14/01/2026

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