

# Felimazole 5 mg/ml oral solution for cats

Authorised

- Thiamazole

## Product identification

**Medicine name:**

Felimazole 5 mg/ml oral solution for cats

Felimazole 5 mg/ml soluzione orale per gatti

**Active substance:**

Thiamazole

**Target species:**

Cat

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Thiamazole

5.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**

Oral solution

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

QH03BB02

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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:**

Italy

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**Package description:**

Polyethylene terephthalate (PET) amber bottle of 100 ml, closed with a low density polyethylene (LDPE) plug and a high density polyethylene (HDPE) closure. The veterinary medicinal product is supplied with a 1 ml polyethylene (PE) / polypropylene (PP) measuring syringe for administration of the solution to the animal. The syringe is graduated in 0.25 mg increments up to 5 mg. Each closed bottle and accompanying syringe is contained in a cardboard carton.

Polyethylene terephthalate (PET) amber bottle of 30 ml, closed with a low density polyethylene (LDPE) plug and a high density polyethylene (HDPE) closure. The veterinary medicinal product is supplied with a 1 ml polyethylene (PE) / polypropylene (PP) measuring syringe for administration of the solution to the animal. The syringe is graduated in 0.25 mg increments up to 5 mg. Each closed bottle and accompanying syringe is contained in a cardboard carton.

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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 8 of Regulation (EU) 2019/6)

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**Marketing authorisation holder:**

Dechra Regulatory B.V.

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**Marketing authorisation date:**

9/01/2024

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**Manufacturing sites for batch release:**

Genera d.d.

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

103685

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**Date of authorisation status change:**

9/01/2024

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0505/004

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Czechia Denmark Finland France  
Germany Greece Hungary Italy 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

Combined File of all Documents

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

Summary of Product Characteristics