

# Milbeguard Duo 12.5 mg / 125 mg chewable tablets for dogs

Authorised

- Milbemyacin oxime
- Praziquantel

## Product identification

**Medicine name:**

Milbeguard Duo 12.5 mg / 125 mg chewable tablets for dogs

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**Active substance:**

Milbemyacin oxime

Praziquantel

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Milbemyacin oxime

12.50 milligram(s) / 1.00 Tablet

Praziquantel

125.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Chewable 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 except for some pack sizes

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**Authorisation status:**

Valid

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**Authorised in:**

France

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**Package description:**

Polyamide-Aluminium-Polyvinyl chloride / aluminium heat sealed blisters: Cardboard box with 1 blister of 2 tablets (2 tablets).

Polyamide-Aluminium-Polyvinyl chloride / aluminium heat sealed blisters: Cardboard box with 2 blisters of 2 tablets (4 tablets).

Polyamide-Aluminium-Polyvinyl chloride / aluminium heat sealed blisters: Cardboard box with 5 blisters of 2 tablets (10 tablets).

Polyamide-Aluminium-Polyvinyl chloride / aluminium heat sealed blisters: Cardboard box with 12 blisters of 2 tablets (24 tablets).

Polyamide-Aluminium-Polyvinyl chloride / aluminium heat sealed blisters: Cardboard box with 24 blisters of 2 tablets (48 tablets).

Polyamide-Aluminium-Polyvinyl chloride / aluminium heat sealed blisters: Cardboard box with 50 blisters of 2 tablets (100 tablets).

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 18 of Regulation (EU) 2019/6)

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**Marketing authorisation holder:**

Ceva Sante Animale

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**Marketing authorisation date:**

12/09/2023

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**Manufacturing sites for batch release:**

Ceva Sante Animale

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/2907989 8/2023

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**Date of authorisation status change:**

12/09/2023

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0780/002

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
France Germany Greece Hungary Italy Latvia Lithuania Luxembourg  
Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain  
Sweden United Kingdom (Northern Ireland)

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**Generic of:**

600000042342

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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

English (PDF)

Published on: 5/04/2026

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### Package Leaflet and Labelling

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

### Combined File of all Documents