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# MILBETAB 2.5 MG/25 MG CHEWABLE TABLETS FOR SMALL DOGS AND PUPPIES

Authorised

- Milbemycin oxime
- Praziquantel

## Product identification

**Medicine name:**
MILBETAB 2.5 MG/25 MG CHEWABLE TABLETS FOR SMALL DOGS AND PUPPIES

**Active substance:**
Milbemycin oxime

Praziquantel

**Target species:**
Dog

**Route of administration:**
Oral use

## Product details

**Active substance and strength:**
Milbemycin oxime
2.50 milligram(s) / 1.00 Tablet

Praziquantel
25.00 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**
Chewable tablet

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QP54AB51

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

**Authorisation status:**
Valid

**Authorised in:**
Czechia

**Package description:**
Box with 1 blister of 4 chewable tablets
Box with 2 blisters of 5 chewable tablets
Box with 3 blisters of 4 chewable tablets
Box with 5 blisters of 4 chewable tablets
Box with 6 blisters of 4 chewable tablets
Box with 12 blisters of 4 chawable tablets
Box with 1 blister of 2 chewable tablets
Box with 2 blisters of 2 chewable tablets
Box with 1 blister of 5 chewable
Box with 5 blister of 2 chewable tablets
Box with 6 blisters of 2 chewable tablets
Box with 3 blisters of 5 chewable tablets
Box with 10 blister of 2 chewable tablets
Box with 5 blisters of 5 chewable tablets
Box with 6 blisters of 5 chewable tablets
Box with 7 blisters of 5 chewable tablets
Box with 8 blisters of 5 chewable tablets
Box with 9 blisters of 5 chewable tablets
Box with 24 blisters of 2 chewable tablets
Box with 10 blisters of 5 chewable tablets

Box with 12 blisters of 2 chewable tablets

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Generic application (Article 18 of Regulation (EU) 2019/6)

**Marketing authorisation holder:**
Chanelle Pharmaceuticals Manufacturing Limited

**Marketing authorisation date:**
19/11/2025

**Manufacturing sites for batch release:**
Chanelle Pharmaceuticals Manufacturing Limited

**Responsible authority:**
Institute For State Control Of Veterinary Biologicals And Medicaments

**Authorisation number:**
96/056/25-C

**Date of authorisation status change:**
19/11/2025

**Reference member state:**
France

**Procedure number:**
FR/V/0488/002

**Concerned member states:**
Austria Belgium Czechia Finland Germany Hungary Ireland Italy Netherlands Norway Poland Portugal Spain Sweden

**Generic of:**

600000043327

To consult adverse reactions on veterinary medicinal products please go to 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.

Package Leaflet

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

Labelling

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

Combined File of all Documents

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