

# Mektix 2.5 mg/25 mg film-coated tablets for small dogs and puppies weighing at least 0.5 kg

Authorised

- Milbemycin oxime
- Praziquantel

## Product identification

**Medicine name:**

Mektix 2.5 mg/25 mg film-coated tablets for small dogs and puppies weighing at least 0.5 kg

---

**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:**

Film-coated tablet

---

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

QP54AB51

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription except for some pack sizes

---

**Authorisation status:**

Valid

---

**Authorised in:**

France

---

**Package description:**

Blister packs consisting of cold formed OPA/Al/PVC foil and aluminium foil. Package sizes:Cardboard box with 1 blister of 2 tablets.

Blister packs consisting of cold formed OPA/Al/PVC foil and aluminium foil. Package sizes:Cardboard box with 1 blister of 4 tablets.

Blister packs consisting of cold formed OPA/Al/PVC foil and aluminium foil. Package sizes:Cardboard box with 12 blisters, each blister contains 4 tablets (total 48 tablets).

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

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

---

**Marketing authorisation holder:**

KRKA tovarna zdravil d.d. Novo mesto

---

**Marketing authorisation date:**

2/09/2021

---

**Manufacturing sites for batch release:**

KRKA tovarna zdravil d.d. Novo mesto  
Krka-Farma d.o.o.

---

**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

---

**Authorisation number:**

FR/V/2875835 0/2021

---

**Date of authorisation status change:**

2/09/2021

---

**Reference member state:**

Ireland

---

**Procedure number:**

IE/V/0526/003

---

**Concerned member states:**

Austria Belgium Cyprus Denmark Finland France Germany Greece Hungary  
Italy Latvia Lithuania Netherlands Norway Portugal Romania Spain Sweden  
United Kingdom (Northern Ireland)

---

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: 12/10/2025

Download

Package Leaflet and Labelling

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

ie-puar-mr-iev0526003-mektix-25-mg25-mg-film-coated-tablets-for-small-do-en.pdf