# 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

**Authorisation status:**
Valid

**Authorised in:**
Romania

**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:**

9/11/2021

**Manufacturing sites for batch release:**
KRKA tovarna zdravil d.d. Novo mesto
Krka-Farma d.o.o.

**Responsible authority:**
Institute For Control Of Biological Products And Veterinary Medicines

**Authorisation number:**
210185

**Date of authorisation status change:**
7/08/2023

**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

## Documents

Summary of Product Characteristics

English (PDF)
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