

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

Authorised

- Milbemyacin 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

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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

2.50 milligram(s) / 1.00 Tablet

Praziquantel

25.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Film-coated 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

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**Authorisation status:**

Valid

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**Authorised in:**

Denmark

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**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).

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

KRKA tovarna zdravil d.d. Novo mesto

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**Marketing authorisation date:**

29/09/2022

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**Manufacturing sites for batch release:**

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

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**Responsible authority:**

Danish Medicines Agency

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**Authorisation number:**

64907

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**Date of authorisation status change:**

29/09/2022

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0526/003

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**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)

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To consult adverse reactions on veterinary medicinal products please go to  
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## Documents

Summary of Product Characteristics

English (PDF)

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