

# Mektix 4 mg/10 mg chewable tablets for small cats and kittens weighing at least 0.5 kg

Authorised

- Milbemycin oxime
- Praziquantel

## Product identification

**Medicine name:**

Mektix 4 mg/10 mg chewable tablets for small cats and kittens weighing at least 0.5 kg

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**Active substance:**

Milbemycin oxime  
Praziquantel

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**Target species:**

Cat

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Milbemycin oxime

4.00 milligram(s) / 1.00 Tablet

Praziquantel

10.00 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 not subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Netherlands

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**Package description:**

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

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

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

29/04/2019

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**Manufacturing sites for batch release:**

KRKA tovarna zdravil d.d. Novo mesto  
Krka-Farma d.o.o.  
TAD Pharma GmbH

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 124200

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**Date of authorisation status change:**

27/01/2022

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0527/001

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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  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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

Summary of Product Characteristics

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