

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

Active substance:

Milbemycin oxime
Praziquantel

Target species:

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Milbemycin oxime

4.00 milligram(s) / 1.00 Tablet

Praziquantel

10.00 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:

Greece

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.

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:

23/04/2025

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Krka-Farma d.o.o.

TAD Pharma GmbH

Responsible authority:

National Organization For Medicines

Authorisation number:

47108/24-04-2025/K-0239103

Date of authorisation status change:

21/03/2022

Reference member state:

Ireland

Procedure number:

IE/V/0527/001

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)

Published on: 30/06/2024

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Package Leaflet

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