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Mektix 12.5 mg/125 mg film-coated tablets for dogs weighing at least 5 kg

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:
Mektix 12.5 mg/125 mg film-coated tablets for dogs weighing at least 5 kg

Active substance:
Milbemycin oxime

Praziquantel

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Milbemycin oxime
12.50 milligram(s) / 1.00 Tablet

Praziquantel
125.00 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:
Sweden

Available in:
Sweden

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:
5/01/2022

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

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
61019

Date of authorisation status change:
5/01/2022

Reference member state:
Ireland

Procedure number:
IE/V/0526/004

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
Updated on: 13/03/2026
Download

Package Leaflet

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Labelling

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

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