

Mektix 12.5 mg/125 mg chewable tablets for dogs weighing at least 5 kg

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:

Mektix 12.5 mg/125 mg chewable 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:

Chewable 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:

France

Package description:

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

29/04/2019

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto
TAD Pharma GmbH

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/3804125 9/2019

Date of authorisation status change:

29/04/2019

Reference member state:

Ireland

Procedure number:

IE/V/0526/002

Concerned member states:

Austria Belgium Cyprus Denmark Finland France Germany Greece Hungary
Italy Latvia Lithuania Netherlands 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

Package Leaflet and Labelling

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