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
Mektix vet 4 mg/10 mg Filmdragerad tablett

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 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. 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:
26/08/2019

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

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
58597

Date of authorisation status change:
26/08/2019

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: 11/02/2022
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Package Leaflet

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Labelling

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