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Amcofen 16 mg/40 mg chewable tablets for cats weighing at least 2 kg

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:
Amcofen 16 mg/40 mg chewable tablets for cats weighing at least 2 kg

Active substance:
Milbemycin oxime

Praziquantel

Target species:
Cat

Route of administration:
Oral use

Product details

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

Praziquantel
40.00 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:
Poland

Available in:
Poland

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:
4/07/2019

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

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2891

Date of authorisation status change:
4/07/2019

Reference member state:
Ireland

Procedure number:
IE/V/0525/002

Concerned member states:
Belgium Bulgaria Croatia Czechia Estonia France Germany Hungary Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Labelling

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

Package Leaflet

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

Summary of Product Characteristics

English (PDF)
Published on: 26/08/2024
Updated on: 13/03/2026
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