

Amcofen 2.5 mg/25 mg chewable tablets for small dogs and puppies weighing at least 0.5 kg

Authorised

- Milbemyacin oxime
- Praziquantel

Product identification

Medicine name:

Amcofen 2.5 mg/25 mg chewable tablets for small dogs and puppies weighing at least 0.5 kg

Active substance:

Milbemyacin oxime
Praziquantel

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Milbemyacin oxime

2.50 milligram(s) / 1.00 Tablet

Praziquantel

25.00 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:

Hungary

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:

29/03/2019

Manufacturing sites for batch release:

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

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

29/03/2019

Reference member state:

Ireland

Procedure number:

IE/V/0524/001

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

Summary of Product Characteristics

English (PDF)

Published on: 3/05/2024

Download