

Milbenin 12.5 mg/125 mg chewable tablets for dogs

Authorised

- Milbemyacin oxime
- Praziquantel

Product identification

Medicine name:

Milbenin 12.5 mg/125 mg chewable tablets for dogs

Active substance:

Milbemyacin oxime

Praziquantel

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Milbemyacin 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:

2 or 4 tablets in OPA-Al-PVC/AL blisters. 12 blisters of 4 chewable tablets introduced in carton box.

2 or 4 tablets in OPA-Al-PVC/AL blisters. 1 blister of 4 chewable tablets introduced in carton box.

2 or 4 tablets in OPA-Al-PVC/AL blisters. 1 blister of 2 chewable tablets introduced in carton box.

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:

Pharma VIM Kft.

Marketing authorisation date:

27/03/2023

Manufacturing sites for batch release:

Vim Spectrum S.R.L.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/2098422 3/2023

Date of authorisation status change:

27/03/2023

Reference member state:

Hungary

Procedure number:

HU/V/0146/002

Concerned member states:

Belgium France Germany Greece Italy Luxembourg Netherlands Portugal
Romania Spain

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 15/11/2023

Updated on: 14/03/2026

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Package Leaflet and Labelling

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Combined File of all Documents