

MILBETAB 12,5 MG/125 MG CHEWABLE TABLETS FOR DOGS

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:

MILBETAB 12,5 MG/125 MG CHEWABLE TABLETS FOR DOGS

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

Authorisation status:

Valid

Authorised in:

Germany

Package description:

Box with 12 blister strips of 4 chewable tablets
Box with 6 blisters strips of 4 chewable tablets
Box with 5 blister strips of 4 chewable tablets
Box with of 3 blisters strips of 4 chewable tablets
Box with 2 blister strip of of 5 chewable tablets
Box with 1 blister strip of 4 chewable tablets
Box with 10 blister strips of 5 chewable tablets
Box with 24 blisters strips of 2 chewable tablets
Box with 9 blister strips of 5 chewable tablets
Box with 8 blister strips of 5 chewable tablets
Box with 7 blister strips of 5 chewable tablets
Box with 6 blister strips of 5 chewable tablets
Box with 5 blister strips of 5 chewable tablets,
Box with 12 blister strips of 2 chewable tablets
Box with 10 blisters strips of 2 chewable tablets
Box with 3 blister strips of 5 chewable tablets
Box with of 6 blister strips of 2 chewable tablets
Box with 5 blisters strips of 2 chewable tablets
Box with 1 blister strips of 5 chewable tablets
Box with 2 blister strips of 2 chewable tablets
1 box with 1 blister of 2 chewable tablets

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:

Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:

15/10/2025

Manufacturing sites for batch release:

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

V7016738.00.00

Date of authorisation status change:

15/10/2025

Reference member state:

France

Procedure number:

FR/V/0488/001

Concerned member states:

Austria Belgium Czechia Finland Germany Hungary Ireland Italy
Netherlands Norway Poland Portugal Spain Sweden

Generic of:

600000043292

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

Documents

Combined File of all Documents

English (PDF)

Published on: 13/03/2026

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