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MILBETAB 2.5 MG/25 MG CHEWABLE TABLETS FOR SMALL DOGS AND PUPPIES

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:
MILBETAB 2.5 MG/25 MG CHEWABLE TABLETS FOR SMALL DOGS AND PUPPIES

Active substance:
Milbemycin oxime

Praziquantel

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Milbemycin 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:
Czechia

Package description:
Box with 1 blister of 4 chewable tablets
Box with 2 blisters of 5 chewable tablets
Box with 3 blisters of 4 chewable tablets
Box with 5 blisters of 4 chewable tablets
Box with 6 blisters of 4 chewable tablets
Box with 12 blisters of 4 chawable tablets
Box with 1 blister of 2 chewable tablets
Box with 2 blisters of 2 chewable tablets
Box with 1 blister of 5 chewable
Box with 5 blister of 2 chewable tablets
Box with 6 blisters of 2 chewable tablets
Box with 3 blisters of 5 chewable tablets
Box with 10 blister of 2 chewable tablets
Box with 5 blisters of 5 chewable tablets
Box with 6 blisters of 5 chewable tablets
Box with 7 blisters of 5 chewable tablets
Box with 8 blisters of 5 chewable tablets
Box with 9 blisters of 5 chewable tablets
Box with 24 blisters of 2 chewable tablets
Box with 10 blisters of 5 chewable tablets

Box with 12 blisters 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:
19/11/2025

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/056/25-C

Date of authorisation status change:
19/11/2025

Reference member state:
France

Procedure number:
FR/V/0488/002

Concerned member states:
Austria Belgium Czechia Finland Germany Hungary Ireland Italy Netherlands Norway Poland Portugal Spain Sweden

Generic of:

600000043327

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

Documents

Summary of Product Characteristics

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.

Labelling

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

Combined File of all Documents

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