# Milbepar 25 mg / 250 mg chewable tablets for large dogs

Authorised

- Milbemycin oxime
- Praziquantel

## Product identification

**Medicine name:**
Milbepar 25 mg / 250 mg chewable tablets for large dogs

**Active substance:**
Milbemycin oxime

Praziquantel

**Target species:**
Dog

**Route of administration:**
Oral use

## Product details

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

Praziquantel

250.00 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:**
Ireland

**Package description:**
Polyamide-Aluminium-Polyvinyl chloride / aluminium heat sealed blisters: Cardboard box with 1 blister of 2 tablets (2 tablets).
Polyamide-Aluminium-Polyvinyl chloride / aluminium heat sealed blisters: Cardboard box with 2 blisters of 2 tablets (4 tablets).
Polyamide-Aluminium-Polyvinyl chloride / aluminium heat sealed blisters: Cardboard box with 5 blisters of 2 tablets (10 tablets).
Polyamide-Aluminium-Polyvinyl chloride / aluminium heat sealed blisters: Cardboard box with 12 blisters of 2 tablets (24 tablets).
Polyamide-Aluminium-Polyvinyl chloride / aluminium heat sealed blisters: Cardboard box with 24 blisters of 2 tablets (48 tablets).
Polyamide-Aluminium-Polyvinyl chloride / aluminium heat sealed blisters: Cardboard box with 50 blisters of 2 tablets (100 tablets).

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Hybrid application – change in strength (Article 19(1)(a) of Regulation (EU) 2019/6)

**Marketing authorisation holder:**
Ceva Sante Animale

**Marketing authorisation date:**
12/07/2024

**Manufacturing sites for batch release:**
Ceva Sante Animale

**Responsible authority:**
Health Products Regulatory Authority

**Authorisation number:**
VPA10815/069/003

**Date of authorisation status change:**
12/07/2024

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0887/003

**Concerned member states:**
France

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

## Documents

Summary of Product Characteristics

Combined File of all Documents

ie-puar-mr-iev0887003-milbepar-25-mg--250-mg-chewable-tablets-for-large--en.pdf