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# Milbeguard Duo 12.5 mg / 125 mg chewable tablets for dogs

Authorised

- Milbemycin oxime
- Praziquantel

## Product identification

**Medicine name:**
Milbeguard Duo 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:**
Lithuania

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

**Marketing authorisation holder:**
Ceva Sante Animale

**Marketing authorisation date:**
16/10/2023

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

**Responsible authority:**
State Food And Veterinary Service

**Authorisation number:**
LT/2/23/2780/001-006

**Date of authorisation status change:**
16/10/2023

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0780/002

**Concerned member states:**
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

**Generic of:**
600000042342

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# Documents

Combined File of all Documents

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