

Milbeguard Duo 4 mg / 10 mg film-coated tablets for small cats and kittens

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:

Milbeguard Duo 4 mg / 10 mg film-coated tablets for small cats and kittens

Active substance:

Milbemycin oxime

Praziquantel

Target species:

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Milbemycin oxime

4.00 milligram(s) / 1.00 Tablet

Praziquantel
10.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Film-coated tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB51

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria

Available in:

Austria

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:

14/09/2023

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

841853

Date of authorisation status change:

14/09/2023

Reference member state:

Ireland

Procedure number:

IE/V/0780/004

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:

600000042379

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

Documents

Package Leaflet

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Summary of Product Characteristics

English (PDF)

Published on: 5/04/2026

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Labelling

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