Milbeguard Duo 25 mg / 250 mg chewable tablets for large dogs

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:
Milbeguard Duo 25 mg / 250 mg chewable tablets for large dogs
Milbeguard duo 25 mg/250 mg comprimés à croquer pour chiens
Milbeguard Duo 25 mg/250 mg kauwtabletten voor grote honden
Milbeguard Duo 25 mg/250 mg Kautabletten für große Hunde

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

Withdrawal period by route of administration:
Oral use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AB51

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
Polyamide-Aluminium-Polyvinyl chloride / aluminium heat sealed blisters: Cardboard box with 50 blisters of 2 tablets (100 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 12 blisters of 2 tablets (24 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 2 blisters of 2 tablets (4 tablets).
Polyamide-Aluminium-Polyvinyl chloride / aluminium heat sealed blisters: Cardboard box with 1 blister of 2 tablets (2 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:
13/11/2023

Manufacturing sites for batch release:
CEVA Santé Animale

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V661948

Date of authorisation status change:
13/11/2023

Reference member state:
Ireland

Procedure number:
IE/V/0780/003

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)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 3/05/2024
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Labelling

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Package Leaflet

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