Milbeguard Duo 2.5 mg / 25 mg chewable tablets for small dogs and puppies

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:
Milbeguard Duo 2.5 mg / 25 mg chewable tablets for small dogs and puppies
MILBEGUARD DUO 2,5 MG/25 MG COMPRIMES A CROQUER POUR PETITS CHIENS ET CHIOTS

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

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 except for some pack sizes

Authorisation status:
Valid

Authorised in:
France

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

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
12/09/2023

Manufacturing sites for batch release:
CEVA Santé Animale

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/3667790 8/2023

Date of authorisation status change:
12/09/2023

Reference member state:
Ireland

Procedure number:
IE/V/0780/001

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
