

Milbetab 12.5 mg/125 mg Tablets for Dogs

Authorised

- Milbemyacin oxime
- Praziquantel

Product identification

Medicine name:

Milbetab 12.5 mg/125 mg Tablets for Dogs

Active substance:

Milbemyacin oxime

Praziquantel

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Milbemyacin oxime

12.50 milligram(s) / 1.00 Tablet

Praziquantel

125.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

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:

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Cardboard box with 2 (1 blister strip of 2)

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Cardboard box with 4 (1 blister strip of 4)

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Cardboard box with 4 (2 blister strips of 2)

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Cardboard box with 8 (1 blister strip of 8)

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Cardboard box with 8 (2 blister strips of 4)

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Cardboard box with 8 (4 blister strips of 2)

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Cardboard box with 10 (1 blister strip of 10)

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Cardboard box with 20 (10 blister strips of 2)

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Cardboard box with 20 (2 blister strips of 10)

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Cardboard box with 30 (3 blister strips of 10)

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Cardboard box with 50 (5 blister strips of 10)

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Cardboard box with 100 (10 blister strips of 10)

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Cardboard box with 200 (20 blister strips of 10)

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Cardboard box with 500 tablets (50 blister strips of 10).

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Multipack of 10 individual packs of 2 tablets.

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Multipack of 10 individual packs of 20 tablets.

Blister packs made up of a cold form laminate of OPA/ALU/PVC and a 20µm hard temper aluminium foil. Multipack of 10 individual packs of 50 tablets.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:

28/08/2023

Manufacturing sites for batch release:

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

28/08/2023

Reference member state:

Ireland

Procedure number:

IE/V/0399/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland
France Germany Greece Hungary Italy Latvia Lithuania Luxembourg
Netherlands Norway Poland Portugal Slovakia Spain Sweden

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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

Combined File of all Documents