

Milbetab 12.5 mg/125 mg Tablets for Dogs

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:

Milbetab 12.5 mg/125 mg 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:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB51

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Latvia

Package description:

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

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:

2/08/2019

Manufacturing sites for batch release:

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/DCP/19/0045

Date of authorisation status change:

2/08/2019

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 Luxembourg Netherlands
Norway Poland Portugal Slovakia Spain Sweden

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

Documents

Combined File of all Documents

Package Leaflet

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

Summary of Product Characteristics

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.