# MILBEVET 2.5 MG/25 MG CHEWABLE TABLETS FOR SMALL DOGS AND PUPPIES

Authorised

- Milbemycin oxime
- Praziquantel

# Product identification

# Medicine name:

MILBEVET 2.5 MG/25 MG CHEWABLE TABLETS FOR SMALL DOGS AND PUPPIES MILBEMAX CHEW 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 TabletPraziquantel25.00 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:

Box of 1 Aluminium/aluminium blisters (OPA/Al/PVC/Al/Heat-Seal coating) of 2 tablets box of 12 aluminium strips (polyester/Al/PE) of 4 tablets box of 1 aluminium strips (polyester/Al/PE) of 4 tablets box of 1 aluminium strips (polyester/Al/PE) of 2 tablets Box of 12 Aluminium/aluminium blisters (OPA/Al/PVC/Al/Heat-Seal coating) of 4 tablets Box of 1 Aluminium/aluminium blisters (OPA/Al/PVC/Al/Heat-Seal coating) of 4 tablets

# Additional information

### **Entitlement type:**

Marketing Authorisation

# Legal basis of product authorisation:

Informed consent application (Article 13c of Directive No 2001/82/EC)

### Marketing authorisation holder:

Elanco GmbH

### Marketing authorisation date:

7/09/2020

### Manufacturing sites for batch release:

Elanco France S.A.S

# **Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

### Authorisation number:

FR/V/3899513 6/2020

### Date of authorisation status change:

7/09/2020

### **Reference member state:**

France

### **Procedure number:**

FR/V/0434/004

### **Concerned member states:**

Netherlands United Kingdom (Northern Ireland)

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

# Documents

Summary of Product Characteristics

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

Package Leaflet and Labelling

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

Combined File of all Documents

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