

MILBEVET 12.5 MG/125 MG CHEWABLE TABLETS FOR DOGS

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:

MILBEVET 12.5 MG/125 MG CHEWABLE 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:

Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB51

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Available in:

Netherlands

Package description:

Box of 1 Aluminium/aluminium blisters (OPA/Al/PVC/Al/Heat-Seal coating) of 2 tablets

Box of 24 Aluminium/aluminium blisters (OPA/Al/PVC/Al/Heat-Seal coating) of 4 tablets

Box of 12 aluminium strips (polyester/Al/PE) of 4 tablets

Box of 12 Aluminium/aluminium blisters (OPA/Al/PVC/Al/Heat-Seal coating) of 4 tablets

Box of 1 aluminium strips (polyester/Al/PE) of 4 tablets

Box of 1 Aluminium/aluminium blisters (OPA/Al/PVC/Al/Heat-Seal coating) of 4 tablets

Box of 1 aluminium strips (polyester/Al/PE) of 2 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:

4/05/2021

Manufacturing sites for batch release:

Elanco France S.A.S

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 127659

Date of authorisation status change:

6/04/2022

Reference member state:

France

Procedure number:

FR/V/0434/003

Concerned member states:

Netherlands United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)

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