

MILPRO 12.5 MG/125 MG FILM-COATED TABLETS FOR DOGS

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:

MILPRO 12.5 MG/125 MG FILM-COATED TABLETS FOR DOGS

Milpro 12,5 mg/125 mg comprimidos revestidos para cães

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:

Film-coated 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

Authorisation status:

Valid

Authorised in:

Portugal

Available in:

Portugal

Package description:

Cardboard box of 2 tablets containing 1 blister of 2 tablets

Cardboard box of 48 tablets containing 24 blisters of 2 tablets

Cardboard box of 24 tablets containing 12 blisters of 2 tablets

Cardboard box of 4 tablets containing 2 blisters of 2 tablets

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Virbac

Marketing authorisation date:

2/04/2014

Manufacturing sites for batch release:

Virbac

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

785/02/14DFVPT

Date of authorisation status change:

19/03/2025

Reference member state:

France

Procedure number:

FR/V/0390/002

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland
Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg
Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain
Sweden United Kingdom (Northern Ireland)

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www.adrreports.eu/vet

Documents

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