

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 õhukese polümeerikattega tabletid koertele

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB51

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Estonia

Available in:

Estonia

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:

13/04/2014

Manufacturing sites for batch release:

Virbac

Responsible authority:

State Agency Of Medicines

Authorisation number:

1832

Date of authorisation status change:

13/04/2014

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)

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

Documents

Summary of Product Characteristics

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