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 δισκία επικαλυμμένα με λεπτό υμένιο για σκύλους

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:
Cyprus

Available in:
Cyprus

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:
7/05/2014

Manufacturing sites for batch release:
Virbac

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00449V

Date of authorisation status change:
8/05/2019

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

Combined File of all Documents

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