

Milprazon 2.5 mg/25 mg tablets for small dogs and puppies weighing at least 0.5 kg

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:

Milprazon 2.5 mg/25 mg tablets for small dogs and puppies weighing at least 0.5 kg
MILPRAZON 2,5 mg/25 mg comprimate pentru câini de talie mică și căței cu greutate de cel puțin 0,5 kg

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 Tablet

Praziquantel

25.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB51

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Romania

Available in:

Romania

Package description:

Blister packs consisting of cold formed OPA/Al/PVC foil and aluminium foil. Box with 1 blister of 2 tablets.

Blister packs consisting of cold formed OPA/Al/PVC foil and aluminium foil. Box with 1 blister of 4 tablets.

Blister packs consisting of cold formed OPA/Al/PVC foil and aluminium foil. Box with 12 blisters, each blister contains 4 tablets.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:

1/02/2015

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Krka-Farma d.o.o.

TAD Pharma GmbH

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

200077

Date of authorisation status change:

9/04/2024

Reference member state:

Ireland

Procedure number:

IE/V/0462/001

Concerned member states:

Belgium Bulgaria Croatia Cyprus Czechia Estonia France Germany Greece

Hungary Italy Latvia Lithuania Netherlands 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

English (PDF)

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