Milprazon 16 mg/40 mg film-coated tablets for cats weighing at least 2 kg

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:
Milprazon 16 mg/40 mg film-coated tablets for cats weighing at least 2 kg
Milprazon 16 mg - 40 mg Filmomhulde tablet
Milprazon 16 mg - 40 mg Comprimé pelliculé
Milprazon 16 mg - 40 mg Filmtablette

Active substance:
Milbemycin oxime
Praziquantel

Target species:
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Milbemycin oxime
16.00 milligram(s) / 1.00 Tablet

Praziquantel
40.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Film-coated tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AB51

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

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:
22/05/2015

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto
TAD Pharma GmbH
Krka-Farma d.o.o.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V474026

Date of authorisation status change:
22/05/2015

Reference member state:
Ireland

Procedure number:
IE/V/0464/002

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)
Published on: 26/08/2024
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Package Leaflet

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Labelling

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