Milprazon 4 mg/10 mg film-coated tablets for small cats and kittens weighing at least 0.5 kg

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:
Milprazon 4 mg/10 mg film-coated tablets for small cats and kittens weighing at least 0.5 kg
MILPRAZON 4 MG/10 MG COMPRIME PELLICULE POUR PETITS CHATS ET CHATONS PESANT AU MOINS 0,5 KG

Active substance:
Milbemycin oxime

Praziquantel

Target species:
Cat

Route of administration:
Oral use

Product details

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

Praziquantel
10.00 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 except for some pack sizes

Authorisation status:
Valid

Authorised in:
France

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:
13/06/2018

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

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/4079890 7/2018

Date of authorisation status change:
20/03/2020

Reference member state:
Ireland

Procedure number:
IE/V/0464/001

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)
Published on: 26/08/2024
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Package Leaflet and Labelling

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