Alpramil 12 mg/30 mg film-coated tablets for cats weighing at least 3 kg

Authorised

- Praziquantel
- Milbemycin oxime

Product identification

Medicine name:
Alpramil 12 mg/30 mg film-coated tablets for cats weighing at least 3 kg

Active substance:
Praziquantel

Milbemycin oxime

Target species:
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Praziquantel
30.00 milligram(s) / 1.00 Tablet

Milbemycin oxime
12.00 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:
Iceland

Package description:
Box with 10 PVC / PE / PVDC - Aluminium blisters each containing 4 tablets.
Box with 10 PVC / PE / PVDC - Aluminium blisters each containing 2 tablets.
Box with 1 PVC/PE/PVDC - Aluminium blister containing 2 tablets.
Box with 1 PVC/PE/PVDC - Aluminium blister containing 1 tablet.
Box with 25 PVC / PE / PVDC - Aluminium blisters each containing 4 tablets.
Box with 25 PVC / PE / PVDC - Aluminium blisters each containing 2 tablets.
Box with 25 PVC / PE / PVDC - Aluminium blisters each containing 1 tablet.
Box with 10 PVC / PE / PVDC - Aluminium blisters each containing 1 tablet.
Box with 1 PVC / PE / PVDC - Aluminium blister containing 4 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:
Alfasan Nederland B.V.

Marketing authorisation date:
21/06/2022

Manufacturing sites for batch release:
Alfasan Nederland B.V.
Lelypharma B.V.

Responsible authority:
Icelandic Medicines Agency

Authorisation number:
IS/2/22/006/02

Date of authorisation status change:
21/06/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0364/002

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg 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

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

English (PDF)
Published on: 14/01/2026
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