Alpramil 12.5 mg/125 mg tablets for dogs weighing at least 5 kg

Authorised

- Praziquantel
- Milbemycin oxime

Product identification

Medicine name:
Alpramil 12.5 mg/125 mg tablets for dogs weighing at least 5 kg

Active substance:
Praziquantel

Milbemycin oxime

Target species:
Dog

Route of administration:
Oral use

Product details

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

Milbemycin oxime

12.50 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:
Czechia

Package description:
Box with 10 OPA/Aluminium/PVC-Aluminium blisters each containing 1 tablet.
Box with 1 OPA/Aluminium/PVC-Aluminium blister containing 4 tablets.
Box with 25 OPA/Aluminium/PVC-Aluminium blisters each containing 4 tablets.
Box with 25 OPA/Aluminium/PVC-Aluminium blisters each containing 2 tablets.
Box with 25 OPA/Aluminium/PVC-Aluminium blisters each containing 1 tablet.
Box with 10 OPA/Aluminium/PVC-Aluminium blisters each containing 4 tablets.
Box with 10 OPA/Aluminium/PVC-Aluminium blisters each containing 2 tablets.
Box with 1 OPA/Aluminium/PVC-Aluminium blister containing 2 tablets
Box with 1 OPA/Aluminium/PVC-Aluminium blister containing 1 tablet

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:
Alfasan Nederland B.V.

Marketing authorisation date:
3/08/2022

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

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/026/22-C

Date of authorisation status change:
3/08/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0364/005

Concerned member states:
Belgium Bulgaria Croatia Cyprus Czechia Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain

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

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.

Labelling

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

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