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Fatrobendan 1.25 mg, chewable tablets for dogs

Authorised

- Pimobendan

Product identification

Medicine name:
Fatrobendan 1.25 mg, chewable tablets for dogs

Active substance:
Pimobendan

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Pimobendan
1.25 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC01CE90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Greece

Available in:
Greece

Package description:
PVC/PE/PVdC/PE/PVC blister sealed with thermoheated aluminium foil containing 10 tablets.Pack-sizes:- Cardboard box containing 10 blisters of 10 tablets (100 tablets)
PVC/PE/PVdC/PE/PVC blister sealed with thermoheated aluminium foil containing 10 tablets.Pack-sizes:- Cardboard box containing 1 blister of 10 tablets (10 tablets)
PVC/PE/PVdC/PE/PVC blister sealed with thermoheated aluminium foil containing 10 tablets.Pack-sizes:- Cardboard box containing 5 blisters of 10 tablets (50 tablets)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application – change in strength (Article 19(1)(a) of Regulation (EU) 2019/6)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
11/12/2024

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
National Organization For Medicines

Authorisation number:
145051/12-12-2024/K-0261801

Date of authorisation status change:
11/12/2024

Reference member state:
Ireland

Procedure number:
IE/V/0571/001

Concerned member states:
Cyprus Czechia Greece Lithuania Poland Portugal Slovakia Spain 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: 29/09/2024
Updated on: 13/03/2026
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Package Leaflet

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