

Proin 50 mg chewable tablets for dogs

Authorised

- Phenylpropanolamine hydrochloride

Product identification

Medicine name:

Proin 50 mg chewable tablets for dogs

Active substance:

Phenylpropanolamine hydrochloride

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Phenylpropanolamine hydrochloride
50.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG04BX91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Package description:

White high density polyethylene bottle containing a 5 gram desiccant pack and cotton, sealed with a child resistant, foil lined heat sealed white polypropylene cap. Pack size: 60 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:

Pegasus Laboratories Ireland Limited

Marketing authorisation date:

20/03/2019

Manufacturing sites for batch release:

Tairgi Tread-Lia Baile Na Sceilge Teoranta

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

3762 ESP

Date of authorisation status change:

1/01/2023

Reference member state:

Ireland

Procedure number:

IE/V/0361/002

Concerned member states:

Belgium France Germany Italy Netherlands 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: 3/05/2024

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Package Leaflet

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Labelling

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