Phenoleptil 12.5 mg tablets for dogs

Authorised

- Phenobarbital

Product identification

Medicine name:
Phenoleptil 12.5 mg tablets for dogs

Active substance:
Phenobarbital

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Phenobarbital
12.50 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN03AA02

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

Authorisation status:
Valid

Authorised in:
Sweden

Available in:
Sweden

Package description:
Aluminum/PVC strips with 10 tablets packed in cardboard boxes with 5 strips
Aluminum/PVC strips with 10 tablets packed in cardboard boxes with 10 strips
Aluminum/PVC strips with 10 tablets packed in cardboard boxes with 25 strips
Aluminum/PVC strips with 10 tablets packed in cardboard boxes with 50 strips
Aluminum/PVC strips with 10 tablets packed in cardboard boxes with 100 strips
Aluminum/PVC strips with 10 tablets packed in cardboard boxes with 100 strips
Aluminum/PVC strips with 10 tablets packed in cardboard boxes with 100 strips
Aluminum/PVC strips with 10 tablets packed in cardboard boxes with 100 strips
Aluminum/PVC strips with 10 tablets packed in cardboard boxes with 100 strips
Aluminum/PVC strips with 10 tablets packed in cardboard boxes with 100 strips

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:
Dechra Regulatory B.V.

Marketing authorisation date:

12/07/2012

Manufacturing sites for batch release:
Lelypharma B.V.
Genera d.d.

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
45347

Date of authorisation status change:
12/07/2012

Reference member state:
Ireland

Procedure number:
IE/V/0635/001

Concerned member states:
Austria Belgium Czechia Denmark Finland France Germany Greece Hungary Iceland Italy Luxembourg Norway Poland Portugal Slovakia Spain Sweden United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Package Leaflet

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

Summary of Product Characteristics

English (PDF)
Published on: 6/07/2025
Download

Labelling

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

Combined File of all Documents