Phenoleptil 12.5 mg tablets for dogs

Authorised

- Phenobarbital

Product identification

Medicine name:
Phenoleptil 12.5 mg tablets for dogs
Phenoleptil vet 12,5 mg tabletti

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:
Finland

Package description:
Aluminium/PVC/PE/PVdC strips with 10 tablets packed in cardboard boxes with 5 strips
Aluminium/PVC/PE/PVdC strips with 10 tablets packed in cardboard boxes with 10 strips
Aluminium/PVC/PE/PVdC strips with 10 tablets packed in cardboard boxes with 25 strips
Aluminium/PVC/PE/PVdC strips with 10 tablets packed in cardboard boxes with 50 strips
Aluminium/PVC/PE/PVdC 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 50 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 10 strips
Aluminum/PVC strips with 10 tablets packed in cardboard boxes with 5 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:
13/11/2011

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

Responsible authority:
Finnish Medicines Agency

Authorisation number:
29304

Date of authorisation status change:
13/11/2011

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

Summary of Product Characteristics

English (PDF)

Published on: 6/07/2025
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Combined File of all Documents

Package Leaflet

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