Phenoleptil 50 mg tablets for dogs

Authorised

- Phenobarbital

Product identification

Medicine name:
Phenoleptil 50 mg tablets for dogs

Active substance:
Phenobarbital

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Phenobarbital
50.00 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:
Denmark

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:

12/05/2011

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

Responsible authority:
Danish Medicines Agency

Authorisation number:
47895

Date of authorisation status change:
12/05/2011

Reference member state:
Ireland

Procedure number:
IE/V/0635/002

Concerned member states:
Austria Belgium Czechia Denmark Finland France Germany Greece Hungary Iceland Italy Luxembourg Norway Poland Portugal Slovakia Slovenia 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)

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Combined File of all Documents