Phenocoat 12.5 mg film-coated tablets for dogs

Authorised

- Phenobarbital

Product identification

Medicine name:
Phenocoat 12.5 mg film-coated 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:
Film-coated 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:
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 film-coated tablets. Carton box containing 10 film-coated tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 film-coated tablets. Carton box containing 10 tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 film-coated tablets. Carton box containing 30 film-coated tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 film-coated tablets. Carton box containing 40 film-coated tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 film-coated tablets. Carton box containing 50 film-coated tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 film-coated tablets. Carton box containing 60 film-coated tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 film-coated tablets. Carton box containing 70 film-coated tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 film-coated tablets. Carton box containing 80 film-coated tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 film-coated tablets. Carton box containing 100 film-coated tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 film-coated tablets. Carton box containing 250 film-coated tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 90 film-coated tablets. Carton box containing 90 film-coated tablets.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Bibliographic application (Article 22 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Alfasan Nederland B.V.

Marketing authorisation date:
9/10/2025

Manufacturing sites for batch release:
Alfasan Nederland B.V.

Responsible authority:
Finnish Medicines Agency

Authorisation number:
42981

Date of authorisation status change:
9/10/2025

Reference member state:
Netherlands

Procedure number:
NL/V/0397/002

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania 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

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

Package Leaflet

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

Combined File of all Documents

English (PDF)
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