Phenocoat 25 mg film-coated tablets for dogs

Authorised

- Phenobarbital

Product identification

Medicine name:
Phenocoat 25 mg film-coated tablets for dogs

Active substance:
Phenobarbital

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Phenobarbital
25.00 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:
Estonia

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 10 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:
28/04/2025

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

Responsible authority:
State Agency Of Medicines

Authorisation number:
1198425

Date of authorisation status change:
28/04/2025

Reference member state:
Netherlands

Procedure number:
NL/V/0397/003

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)

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Documents

Summary of Product Characteristics

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Combined File of all Documents

English (PDF)
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