

Phenocoat 50 mg film-coated tablets for dogs

Authorised

- Phenobarbital

Product identification

Medicine name:

Phenocoat 50 mg film-coated 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:

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:

Austria

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:

18/04/2025

Manufacturing sites for batch release:

Alfasan Nederland B.V.

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

842682

Date of authorisation status change:

18/04/2025

Reference member state:

Netherlands

Procedure number:

NL/V/0397/004

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

Package Leaflet

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

Summary of Product Characteristics

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

Labelling

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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