Phenosan 12.5 mg chewable tablets for dogs

Authorised

- Phenobarbital

Product identification

Medicine name:
Phenosan 12.5 mg chewable 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:
Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN03AA02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 chewable tablets. Carton box containing 10 chewable tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 chewable tablets. Carton box containing 20 chewable tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 chewable tablets. Carton box containing 30 chewable tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 chewable tablets. Carton box containing 40 chewable tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 chewable tablets. Carton box containing 50 chewable tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 chewable tablets. Carton box containing 60 chewable tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 chewable tablets. Carton box containing 70 chewable tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 chewable tablets. Carton box containing 80 chewable tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 chewable tablets. Carton box containing 100 chewable tablets.
PVDC/PE/PVC-PVC/Aluminium/Paper blister containing 10 chewable tablets. Carton box containing 250 chewable tablets.
PVC/PE/PVDC-PVC/Aluminium/Paper blister containing 10 chewable tablets. Carton box containing 90 chewable 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:
10/04/2025

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

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 132297

Date of authorisation status change:
18/03/2025

Reference member state:
Netherlands

Procedure number:
NL/V/0396/001

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

Combined File of all Documents

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