

Phenosan 50 mg chewable tablets for dogs

Authorised

- Phenobarbital

Product identification

Medicine name:

Phenosan 50 mg chewable tablets for dogs

Phenosan 50 mg compresse masticabili per cani

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:

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:

Italy

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:

19/05/2025

Manufacturing sites for batch release:

Alfasan Nederland B.V.

Responsible authority:

European Medicines Agency

Authorisation number:

105802

Date of authorisation status change:

19/05/2025

Reference member state:

Netherlands

Procedure number:

NL/V/0396/002

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

Combined File of all Documents

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