DIPHENHYDRAMINE HYDROCHLORIDE- diphenhydramine hydrochloride capsule, liquid filled Prodose, Inc.

Diphenhydramine Hydrochloride Capsules USP 50 mg (Dye-free)

Active ingredient

(in each softgel)

Diphenhydramine Hydrochloride USP 50 mg

Purpose

Nighttime sleep-aid

Uses

for relief of occasional sleeplessness

Warnings

Do not use

- for children under 12 years of age
- with any other product containing diphenhydramine, even one used on skin

Ask a doctor before use if you have

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- trouble urinating due to an enlarged prostate gland

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers

When using this product

avoid alcoholic drinks

Stop use and ask a doctor if

sleeplessness persist continuously for more than 2 weeks. Insomnia may be a symptom of a serious underlying medical illness.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

 adults and children 12 years of age and over: 1 softgel (50 mg) at bedtime if needed, or as directed by a doctor

Inactive ingredients

gelatin, glycerin, polyethylene glycol 400, polyethylene glycol 600, propylene glycol, purified water, sorbitol sorbitan solution.

Bulk Package Label

<u>Diphenhyd</u>	dramine Hydrochloride Capsules USP 50mg (Dye Fr	<u>'ee)</u> IMPRINT
	<u>Each Soft Gelatin Capsule Contains:</u> (Diphenhydramine Hydrochloride USP 50mg)	
BATCH NO. :	NDC NO : 68210-5057-1	
GROSS WT :	MFG. DATE :	
NET WT : QUANTITY :	EXP DATE : SHIPPER NO :	
	<u>WARNING</u> KEEP OUT OF REACH OF CHILDREN	
AVOID HI	STORE BETWEEN 20° - 25°C (68° to 77°F). IGH HUMIDITY AND EXCESSIVE HEAT. PROTECT FROM LIGHT.	
111010	BULK SHIPMENT INTENDED FOR FURTHER PROCESSING ONLY. PPROVED, REPACKED IMMEDIATELY AND LABELED IN STRICT WITHTHE F.D. & C ACT AND REGULATIONS.	CONFORMANCE
MANUFACTURED FO	R:	
PRODOSE INC.,		
2004 ORVILLE DRIVE N		
RONKONKOMA, NEW Y UNITED STATES (USA)		
, ,	68210	

DIPHENHYDRAMINE HYDROCHLORIDE

diphenhydramine hydrochloride capsule, liquid filled

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68210-5057	
Route of Administration	ORAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	50 mg	

Inactive Ingredients				
Ingredient Name	Strength			
POLYETHYLENE GLYCOL 600 (UNII: NL4J9F21N9)				
WATER (UNII: 059QF0KO0R)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)				
SORBITOL (UNII: 506T60A25R)				
GELATIN (UNII: 2G86QN327L)				
GLYCERIN (UNII: PDC6A3C0OX)				

Product Characteristics				
Color	white (Clear transparent)	Score	no score	
Shape	OVAL	Size	13mm	
Flavor		Imprint Code	02	
Contains				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:68210- 5057-1	2 in 1 BOX	07/30/2025		
1		3000 in 1 POUCH; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M010	07/30/2025	

Labeler - Prodose, Inc. (119371190)

Revised: 7/2025 Prodose, Inc.