SLEEP AID- diphenhydramine hcl capsule QUALITY CHOICE (Chain Drug Marketing Association)

Drug Facts

Active ingredient (in each softgel)

Diphenhydramine HCL 50 mg

Purpose

Nighttime sleep-aid

Use

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 persists 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 (1-800-222-1222) 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

Other information

· • •

• store between 15-30°C (59°-86°F)

Inactive ingredients

FD&C blue #1, gelatin, glycerin, lecithin, light mineral oil, mannitol, polyethylene glycol, purified water, sorbitol, sorbitan, white ink

Questions or comments?

Call 1-800-935-2362 Monday-Friday 9AM-5PM EST

Principal Display Panel

†Compare to the Active Ingredient in Unisom® SleepGels®

Sleep-AidMaximum Strength

Nighttime Sleep-Aid

Diphenhydramine HCL 50 mg

Help To Reduce Difficulty Falling Asleep

Softgels

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

†This product is not manufactured or distributed by Sanofi Consumer Healthcare NA, distributor of Unisom® SleepGels®.

Distributed by CDMA, Inc

Novi, MI 48375

Package Label





Contains

SLEEP AID						
diphenhydramine hcl capsu	le					
Product Information						
Product Type	HUMAN OTC	DRUG	Item Code (S	Source)	NDC:83	324-151
Route of Administration	ORAL					
Active Ingradient/Activ	o Moloty					
Active Ingredient/Activ	-					
	redient Nam			Basis of S	-	Strength
DIPHENHYDRAMINE HYDROCH (DIPHENHYDRAMINE - UNII:8GTS8		TC2D6JAD40)		DIPHENHYDRAM HYDROCHLORID		50 mg
luce et a luce d'autre						
Inactive Ingredients						
	-	ent Name			9	Strength
FD&C BLUE NO. 1 (UNII: H3R47	7K3TBD)					
GELATIN (UNII: 2G86QN327L)						
GLYCERIN (UNII: PDC6A3C0OX)						
LIGHT MINERAL OIL (UNII: N6K						
POLYETHYLENE GLYCOL, UNS	SPECIFIED (UNI	I: 3WJQ0SDW1	4)			
SORBITAN (UNII: 6092ICV9RU)						
SORBITOL (UNII: 506T60A25R) LECITHIN, SOYBEAN (UNII: 1DI	5600M62)					
MANNITOL (UNII: 30WL53L36A)	50QDM02)					
Product Characteristic	s					
Color	blue	Score			no score	
Shape	OVAL	Size			13mm	
Flavor		Imprint Co	de		PC5	

Pa	ckaging			
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:83324- 151-32	1 in 1 BOX	07/31/2024	
1		32 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
M	arketing	Information		
M	arketing Marketing Category	nformation Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

Labeler - QUALITY CHOICE (Chain Drug Marketing Association) (011920774)

Revised: 5/2024

QUALITY CHOICE (Chain Drug Marketing Association)