SLEEP AID NIGHTTIME- diphenhydramine hydrochloride capsule, liquid filled TOPCO ASSOCIATES LLC

Sleep Aid Nighttime

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 tranguilizers

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 serious underlying medical illness.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

Overdose warning: 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

Other information

store at 20°-25°C (68°-77°F)

Inactive ingredients

FD&C blue #1, FD&C red #40, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, purified water, sorbitol sorbitan solution, titanium dioxide

Questions?

Call toll free: 1-888-423-0139

TopCare®

health

MAXIMUM STRENGTH

- Fall Asleep Fast
- Safe, Non-Habit Forming

COMPARE TO UNISOM[®] SLEEPGELS[®] ACTIVE INGREDIENT*

DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

*This product is not manufactured or distributed by Chattem, Inc., owner of the registered trademark Unisom[®].

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This TopCare® product is laboratory tested to guarantee its highest quality. Your total satisfaction is guaranteed.

Packaging



SLEEP AID NIGHTTIME

diphenhydramine hydrochloride capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:36800-784
Route of Administration	ORAL		

 Active Ingredient/Active Moiety

 Ingredient Name
 Basis of Strength
 Strength

 DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40)
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 DIPHENHYDRAMINE + UNII:8GTS82S83M)
 50 mg

Inactive Ingredients					
Ingredient Name	Strength				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)					
FD&C RED NO. 40 (UNII: WZB9127XOA)					
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)					
GLYCERIN (UNII: PDC6A3C0OX)					

	GLYCOL, UN	POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)							
POVIDONE K30 (UNII: U725QWY32X)									
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)									
WATER (UNII: 059QF0KO0R)									
SORBITOL (UNII: 506T60A25R)									
SORBITAN (UNII: 6092ICV9RU)									
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)									
Product Characteristics									
Color		BLUE	Score		no score				
Shape		OVAL	Size		13mm				
Flavor			Imprint Code		785				
Contains									
Packaging									
# Item Code		Package Des	cription	Marketing Start Date	Marketing End Date				
1 NDC:36800-784	784- 96 in 1 BOTTLE; Type 0: Not a Combination Product		05/01/2021						
Markoting	Inform	ation							
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Marketing Marketing Category			er or Monograph ion	Marketing Star Date	t Marketing End Date				

Labeler - TOPCO ASSOCIATES LLC (006935977)

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