

SLEEP AID- diphenhydramine hcl tablet
Chain Drug Consortium

Premier Value 44-189

Active ingredient (in each tablet)

Diphenhydramine HCl 25 mg

Purpose

Nighttime sleep-aid

Use

helps to reduce difficulty falling asleep

Warnings

Do not use

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

Ask a doctor before use if you have

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- difficulty in urination due to enlargement of the prostate gland

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

- avoid alcoholic beverages
- drowsiness will occur
- use caution when driving a motor vehicle or operating machinery

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 and over: take 2 tablets at bedtime if needed or as directed by a doctor
- children under 12 years: do not use

Other information

- **each tablet contains:** calcium 60 mg
- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- protect from moisture
- see end flap for expiration date and lot number
- store at 25°C (77°F); excursions permitted between 15°C-30°C (59°F-86°F)

Inactive ingredients

croscarmellose sodium, dibasic calcium phosphate dihydrate, FD&C blue #1 aluminum lake, magnesium stearate, microcrystalline cellulose, silicon dioxide, stearic acid

Questions or comments?

1-800-426-9391

Principal Display Panel

**Premier
Value®**

***COMPARE TO THE ACTIVE INGREDIENT IN
ORIGINAL FORMULA SOMINEX®**

Sleep Aid

Diphenhydramine HCl 25 mg

NIGHTTIME SLEEP AID

Wake Up Rested and Refreshed
Doctor preferred sleep ingredient

16 Tablets

PV

INDEPENDENTLY TESTED
SATISFACTION GUARANTEED

**TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER
UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING**

*This product is not manufactured or distributed by Prestige

Brands, Inc., owner of the registered trademark Original Formula Sominex®. 50844 REV0516A18921

Distributed by:
Pharmacy Value Alliance, LLC
407 East Lancaster Avenue,
Wayne, PA 19087

If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.



Sleep Aid

Diphenhydramine HCl 25 mg
NIGHTTIME SLEEP AID



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No print/No varnish
Lot & Exp date

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B-1590-189-21
REV0516A18921

Drug Facts (continued)
store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
Inactive ingredients: croscarmellose sodium, dibasic calcium phosphate dihydrate, FD&C blue #1, aluminum lake, magnesium stearate, microcrystalline cellulose, silicon dioxide.

Drug Facts (continued)
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Questions or comments? 1-800-426-9391

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Other Information
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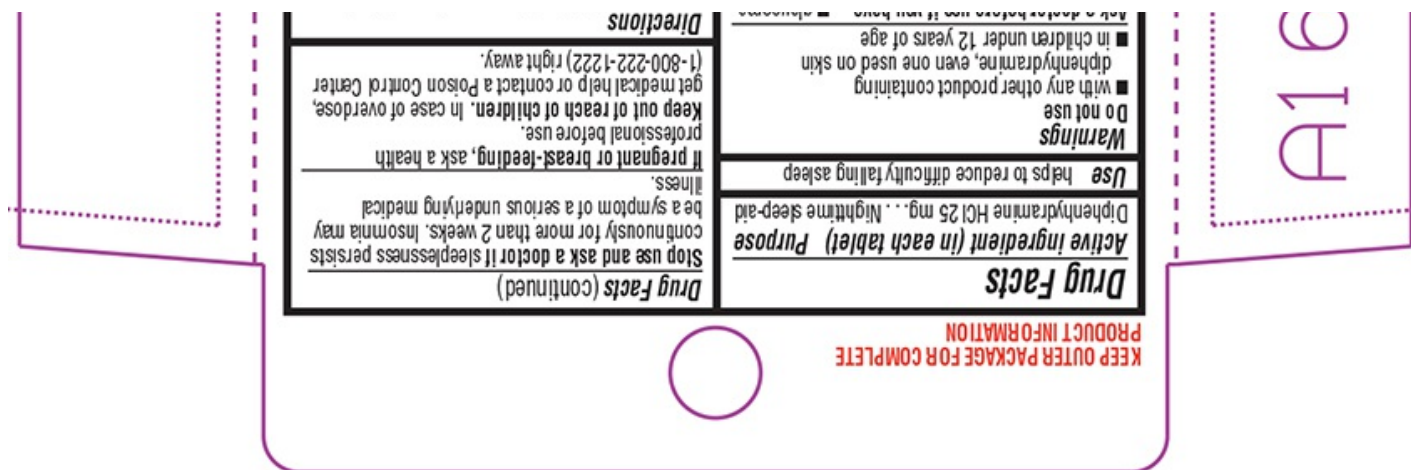
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use caution when driving a motor vehicle or operating machinery

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chronic bronchitis
difficulty in urination due to enlargement of the prostate gland
a breathing problem such as emphysema or chronic bronchitis
glaucoma
if you have

Ask a doctor before use if you have
glaucoma

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Premier Value 44-189

SLEEP AID

diphenhydramine hcl tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-639
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	blue	Score	no score
Shape	ROUND	Size	10mm
Flavor		Imprint Code	44;189
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-639-16	2 in 1 CARTON	04/10/1990	
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:68016-639-32	4 in 1 CARTON	04/10/1990	04/11/2021
2		8 in 1 BLISTER PACK; 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	04/10/1990	

Labeler - Chain Drug Consortium (101668460)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(68016-639)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(68016-639) , pack(68016-639)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(68016-639)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(68016-639)