

**NIGHT TIME SLEEP-AID- diphenhydramine hcl tablet**  
**Advanced Rx LLC**

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**NIGHT TIME SLEEP-AID**

***Drug Facts***

***Active ingredients (in each Caplet)***

Diphenhydramine HCl 25 mg

***Purpose***

Nighttime sleep-aid

***Uses***

- for the relief of occasional sleeplessness
- reduces time to fall asleep if you have difficulty falling asleep

***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
- 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

**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.**

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

***Directions***

- take only one dose (2 Caplets) per day (24 hours)
-

adults & children 12 yrs & over 2 Caplets at bedtime if needed or as directed by a doctor  
children under 12 yrs do not use

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**Other information**

- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- read all product information before using
- TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

**Inactive ingredients**

Microcrystalline Cellulose, Dicalcium Phosphate, Stearic Acid, Croscarmellose Sodium, Colloidal Silicon Dioxide, Hypromellose, Titanium Dioxide, Triacetin, FD&C Red No.40 Lake, FD&C Blue No.2 Aluminum Lake, Polysorbate.

**Questions or comments**

Call toll free 1-800-630-8895

Monday through Friday 9AM - 5PM EST

**Distributed by:**

**Advanced Rx LLC**

1942 NE 163rd St North Miami Beach,  
FL 33162 U.S.A.

**MADE IN THE USA**

**PRINCIPAL DISPLAY PANEL**

**NDC 80513-383-22**

Compare to the active ingredient in VICKS<sup>®</sup> ZzzQuil<sup>®</sup> NIGHTTIME SLEEP-AID\*

**NIGHT TIME SLEEP-AID**

Diphenhydramine HCl 25 mg

220 CAPLETS

\*This product is not manufactured or distributed by The Procter & Gamble Company, owner of the registered trademark VICKS<sup>®</sup> ZzzQuil<sup>®</sup> NIGHTTIME SLEEP-AID



Compare to the active ingredient in VICKS®  
ZzzQuil® NIGHTTIME SLEEP-AID

# NIGHT TIME SLEEP AID

Diphenhydramine HCL 25 mg

**25 MG**

- ✓ Fast Acting
- ✓ Not-Habit Forming

**220**  
CAPLETS  
easy to swallow

### Drug Facts

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### Drug Facts (continued)

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|---------------------------------|---|
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#### Questions or comments

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MADE IN THE USA



Lot No.: 4  
Exp. Date: 80513100093

## NIGHT TIME SLEEP-AID

diphenhydramine hcl tablet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:80513-383
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

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

### Inactive Ingredients

Ingredient Name	Strength
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>ANHYDROUS DIBASIC CALCIUM PHOSPHATE</b> (UNII: L11K75P92J)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>TRIACETIN</b> (UNII: XHX3C3X673)	
<b>FD&amp;C RED NO. 40 ALUMINUM LAKE</b> (UNII: 6T47AS764T)	
<b>FD&amp;C BLUE NO. 2 ALUMINUM LAKE</b> (UNII: 4AQJ3LG584)	

### Product Characteristics

<b>Color</b>	purple	<b>Score</b>	no score
<b>Shape</b>	CAPSULE (Caplet)	<b>Size</b>	11mm
<b>Flavor</b>		<b>Imprint Code</b>	DPZ
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80513-383-22	220 in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2024	12/31/2027

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M010	08/01/2024	12/31/2027

**Labeler** - Advanced Rx LLC (042795108)

Revised: 7/2025

Advanced Rx LLC