

**WAL-SLEEP Z NIGHTTIME- diphenhydramine hcl solution**  
**Walgreen Company**

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**Walgreens 44-002**

***Active ingredient (in each 30 mL dose cup)***

Diphenhydramine HCl 50 mg

***Purpose***

Nighttime sleep-aid

***Uses***

- for 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 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**

- **do not take more than directed**
- mL = milliliter; FL OZ = fluid ounce
- use only enclosed dosing cup designed for use with this product. Do not use any other dosing device.
- take only one dose per day (24 hours)
- adults and children 12 years and over: take 30 mL at bedtime if needed or as directed by a doctor
- children under 12 years: do not use

**Other information**

- **each 30 mL dose cup contains:** sodium 17 mg
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- use by expiration date on package

**Inactive ingredients**

anhydrous citric acid, FD&C blue #1, FD&C red #40, flavor, glycerin, high fructose corn syrup, propylene glycol, purified water, sodium benzoate, sodium chloride, sodium citrate dihydrate, sucralose, sucrose, xanthan gum

**Questions or comments?**

**1-800-426-9391**

**Principal display panel****Walgreens****Compare to Vicks® ZzzQuil® Nighttime Sleep-Aid active ingredient<sup>††</sup>**

NDC 0363-0020-02

**Wal-Sleep Z®****DIPHENHYDRAMINE HCl / NIGHTTIME SLEEP AID**

NIGHTTIME

ALCOHOL FREE

- Non-habit forming

12 FL OZ (355 mL)

NOT FOR TREATING PAIN, COLD OR FLU

BERRY

FLAVOR

**TAMPER EVIDENT: DO NOT USE IF PRINTED**

## NECK WRAP IS BROKEN OR MISSING

††This product is not manufactured or distributed by The Procter & Gamble Company, owner of the registered trademark Vicks® ZzzQuil® Nighttime Sleep-Aid.

50844 REV1120A00202

Walgreens Pharmacist Recommended  
Walgreens Pharmacist Survey

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**Drug Facts (continued)**

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Compare to Vicks® ZzzQuil® Nighttime Sleep-Aid active ingredient††

NDC 0363-0020-02

# Wal-Sleep Z®

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50844 REV1120A00202

No Print / No Varnish Area  
Lot # and Exp. Info



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**ALCOHOL FREE**

• Non-habit forming

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BERRY FLAVOR

ITEM 957126 W00000-0000-0



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ORG0818-F REV1220

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**Walgreens 44-002**

## WAL-SLEEP Z NIGHTTIME

diphenhydramine hcl solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0363-0020
<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	50 mg in 30 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>HIGH FRUCTOSE CORN SYRUP</b> (UNII: XY6UN3QB6S)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>TRISODIUM CITRATE DIHYDRATE</b> (UNII: B22547B95K)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>SUCROSE</b> (UNII: C151H8M554)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	

## Product Characteristics

<b>Color</b>	purple	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	BERRY	<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0020-45	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/12/2016	10/23/2025
2	NDC:0363-0020-02	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/12/2016	10/23/2025
3	NDC:0363-0020-96	2 in 1 PACKAGE	09/12/2016	10/23/2025
3	NDC:0363-0020-02	355 mL in 1 BOTTLE, PLASTIC; 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	09/12/2016	10/23/2025

**Labeler** - Walgreen Company (008965063)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	manufacture(0363-0020) , pack(0363-0020)

Revised: 3/2024

Walgreen Company