

**NIGHTTIME SLEEP AID- diphenhydramine hcl solution**  
**Bi-Mart**

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**Bi-Mart 44-002A**

***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?***

**1-800-935-6737**

***Principal Display Panel******BI-MART***

NDC 37835-319-30

Compare to the active ingredient in  
Vicks® ZzzQuil® NIGHTTIME SLEEP-AID\*

**Nighttime  
Sleep-Aid**

Diphenhydramine HCl 50 mg  
Nighttime Sleep-Aid

- Non-Habit Forming
- Alcohol Free

**Not for treating Cold or Flu**

Berry Flavor

**12 FL OZ (355 mL)**

**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 ORG112000202

BI-MART UNCONDITIONAL MONEY BACK GUARANTEE  
If you are not completely satisfied with your BI-MART  
QUALITY PRODUCT, regardless of the reason, return  
the unused portion and your entire purchase price will  
be cheerfully refunded.

Distributed by:  
BI-MART  
Eugene, OR 97402

**Drug Facts** (continued)

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

Distributed by:  
BI-MART  
Eugene, OR 97402  
L9133-012-11-0

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NDC 37835-319-30

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## NIGHTTIME SLEEP AID

diphenhydramine hcl solution

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37835-319
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40)	DIPHENHYDRAMINE	50 mg

(DIPHENHYDRAMINE - UNII:8GTS82S83M)			HYDROCHLORIDE		in 30 mL
Inactive Ingredients					
Ingredient Name				Strength	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)					
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)					
FD&C RED NO. 40 (UNII: WZB9127XOA)					
GLYCERIN (UNII: PDC6A3C0OX)					
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)					
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)					
WATER (UNII: 059QF0KO0R)					
SODIUM BENZOATE (UNII: OJ245FE5EU)					
SODIUM CHLORIDE (UNII: 451W47IQ8X)					
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)					
SUCRALOSE (UNII: 96K6UQ3ZD4)					
SUCROSE (UNII: C151H8M554)					
XANTHAN GUM (UNII: TTV12P4NEE)					
Product Characteristics					
Color		purple	Score		
Shape			Size		
Flavor		BERRY	Imprint Code		
Contains					
Packaging					
#	Item Code	Package Description		Marketing Start Date	Marketing End Date
1	NDC:37835-319-30	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		04/29/2021	
Marketing Information					
Marketing Category		Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug		M010		04/29/2021	

**Labeler** - Bi-Mart (027630078)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	manufacture(37835-319) , pack(37835-319)