# NIGHTTIME SLEEP AID- diphenhydramine hcl solution L.N.K. International, Inc.

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Sound Body 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?**

1-800-426-9391

### Principal display panel

SOUND**BODY**™

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

NDC 50844-102-00

Alcohol Free

Nighttime Sleep-Aid

Diphenhydramine HCI 50 mg

#### **NIGHTTIME SLEEP-AID**

Non-habit Forming Not for treating Pain, 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 REV1120A00202

Manufactured for Big Lots Stores, Inc.

by LNK INTERNATIONAL, INC.

60 Arkay Drive, Hauppauge, NY 11788 USA V#733000 ITEM#022700202



Sound Body 44-002A

#### NIGHTTIME SLEEP AID

diphenhydramine hcl solution

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50844-102	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				

1	Active Ingredient/Active Moiety			
1	Ingredient Name	Basis of Strength	Strength	
	<b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	50 mg in 30 mL	
1				

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:50844- 102-00	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/04/2020	
2	NDC:50844- 102-02	2 in 1 PACKAGE	03/04/2020	
2		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	03/04/2020		

## Labeler - L.N.K. International, Inc. (038154464)

Establishment				
Name	Address	ID/FEI	Business Operations	
LNK International, Inc.		967626305	manufacture(50844-102) , pack(50844-102)	