

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

Quality Plus 44-002A

Active ingredient (in each 30 mL)

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
- only use the dose cup provided
- 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 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

**Quality
+ Plus**

NDC 50844-200-02

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

**Nighttime
Sleep Aid**

**Diphenhydramine HCl 50 mg
per 30 mL**

Nighttime Sleep-Aid

- Non-habit Forming
- Not For Colds or
For Pain

Berry Flavor

Ages 12 Years and Over

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 ORG022300202

Distributed by
LNK INTERNATIONAL, INC.
60 Arkay Drive
Hauppauge, NY 11788
USA



Quality Plus 44-002A

NIGHTTIME SLEEP AID

diphenhydramine hcl solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50844-200
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
-----------------	-------------------	----------

DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)		DIPHENHYDRAMINE HYDROCHLORIDE	50 mg 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:50844-200-45	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/12/2016	
2	NDC:50844-200-02	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/12/2016	
3	NDC:50844-200-96	2 in 1 PACKAGE	09/12/2016	
3		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	

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

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	manufacture(50844-200) , pack(50844-200)

Revised: 3/2024

L.N.K. International, Inc.