

SLEEP AID NIGHTTIME- diphenhydramine hcl liquid
TARGET Corporation

Drug Facts

Active ingredient (in each 30 mL)

Diphenhydramine HCl 50 mg

Purpose

Nighttime sleep-aid

Use

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

Warnings

Do not use

- in 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
- trouble urinating due to an enlarged 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 (1-800-222-

1222) right away.

Directions

- take only one dose per day (24 hours)
- mL = milliliter
- keep dosing cup with product
- measure only with dosing cup provided. Do not use any other dosing device
- adults and children 12 years and over
 - one dose = 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 24 mg
- store between 20-25°C (68-77°F). Do not refrigerate
- protect from light

Inactive ingredients

citric acid, FD&C blue #1, FD&C red #40, flavor, high fructose corn syrup, polyoxyl 40 stearate, propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium citrate

Questions or comments?

Call **1-800-910-6874**

Principal Display Panel

Compare to active ingredient in ZzzQuil®*

nighttime sleep aid

diphenhydramine HCl 50 mg

non habit-forming

alcohol free

BERRY FLAVOR

FL OZ (mL)

NOT FOR TREATING COLD OR FLU

*This product is not manufactured or distributed by The Procter & Gamble Company. ZzzQuil® is a registered trademark of The Procter & Gamble Company.

Failure to follow these warnings could result in serious consequences

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL AROUND DOSAGE CUP OR UNDER CAP IS BROKEN OR MISSING.

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Minneapolis, MN 55403
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Product Label

Drug Facts (continued)

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or distributed by The Procter &
Gamble Company. ZzzQuil® is a
registered trademark of The
Procter & Gamble Company.

PLD-A373C LB006441



PAPER
BOX



PLASTIC
BOTTLE

Discard Seal,
Empty &
Replace Cap

how2recycle.info

NDC 11673-375-12

Compare to active ingredient
in ZzzQuil®*

**nighttime
sleep aid**

diphenhydramine HCl, 50 mg

non habit-forming
alcohol free

up&up

12 FL OZ (354 mL)

NOT FOR TREATING COLD OR FLU

BERRY
FLAVOR

TARGET Nighttime Sleep Aid

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SLEEP AID NIGHTTIME

diphenhydramine hcl liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-375
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)			

HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)				
POLYOXYL 40 STEARATE (UNII: 13A4J4NH9I)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
SODIUM CITRATE (UNII: 1Q73Q2JULR)				
Product Characteristics				
Color			Score	
Shape			Size	
Flavor		BERRY	Imprint Code	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-375-12	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/31/2016	07/31/2025
2	NDC:11673-375-24	2 in 1 PACKAGE	07/31/2016	07/31/2025
2		354 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	07/31/2016	07/31/2025

Labeler - TARGET Corporation (006961700)