

SLEEP AID NIGHTTIME- diphenhydramine hcl liquid
TARGET Corporation

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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)

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.
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- one dose = 30 mL at bedtime if needed, or as directed by a doctor
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Other information

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PLD-A373C LB006441



NDC 11673-375-12

Compare to active ingredient in ZzzQuil®*

nighttime sleep aid

diphenhydramine HCl, 50 mg

non habit-forming
alcohol free




12 FL OZ (354 mL)

NOT FOR TREATING COLD OR FLU

TARGET Nighttime Sleep Aid

Failure to follow these warnings could result in serious consequences.

Drug Facts

Active ingredient (in each 30 mL)	Purpose
Diphenhydramine HCl 50 mg	Nighttime sleep-aid

Uses

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- reduces time to fall asleep if you have difficulty falling asleep

Warnings

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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: 059QF0K00R)	
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:11673-375-24	2 in 1 PACKAGE	07/31/20 16	
1		354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:11673-375-12	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/31/20 16	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part338	07/31/20 16	

Labeler - TARGET Corporation (006961700)

Revised: 3/2019

TARGET Corporation