

SLEEP AID- 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 23 mg
- store between 20-25°C (68-77°F). Do not refrigerate
- protect from light

Inactive ingredients

citric acid, ethyl alcohol, 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

50 mg Diphenhydramine HCl per 30 mL

- Non habit-forming
- Not for treating cold or flu
- Alcohol 10%

Berry Flavor

FL OZ (mL)

*This product is not manufactured or distributed by The Procter & Gamble Company. Vicks® 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.

Distributed by Target Corporation

Minneapolis, MN 55403

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.
- 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 23 mg
- store between 20-25°C (68-77°F). Do not refrigerate.
- protect from light.

Inactive ingredients citric acid, ethyl alcohol, 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

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

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

NDC 11673-750-12
094 01 0155 R00
C-002262-01-034-0000

Distributed by Target Corporation
Minneapolis, MN 55403
Product of China
TM & ©2024 Target Brands, Inc.

PLD-A262D L8009459

Compare to active ingredient
in Vicks® ZzzQuil®*

Nighttime Sleep Aid

50 mg Diphenhydramine HCl
per 30 mL

- Non-habit forming
- Not for treating cold or flu
- ALCOHOL 10%

12 FL OZ (355 mL)

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

- 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.

TARGET (up&up) Nighttime Sleep Aid Berry Flavor

SLEEP AID

diphenhydramine hcl liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-750
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)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-750-12	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/31/2023	
2	NDC:11673-750-24	2 in 1 PACKAGE	10/31/2023	
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	10/31/2023	

Labeler - TARGET Corporation (006961700)

Revised: 11/2023

TARGET Corporation