ZZZQUIL NIGHTTIME SLEEP-AID- diphenhydramine hydrochloride liquid
The Procter & Gamble Manufacturing Company

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.

ZzzQuil™

NIGHTTIME SLEEP-AID

Drug Facts

Active ingredients
(in each 30 mL dose cup or 2 tablespoons)

Diphenhydramine HCl 50 mg

Purpose

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

- 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 serious underlying medical illness.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

Overdose warning

In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- take only one dose per day (24 hours) - see Overdose warning
- only use dose cup provided

adults & children 12 yrs & over
30 mL at bed time if needed or as directed by a doctor

children under 12 yrs

do not use

Other information

- **each 30 mL dose (2 tablespoons) contains:** sodium 24 mg
- store at controlled room temperature

Inactive ingredients

alcohol, citric acid anhydrous, FD&C Blue No. 1, FD&C Red No. 40, flavor, high fructose corn syrup, polyoxyl 40 stearate, propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium citrate

Questions?

1-877-881-5813

**Dist. by Procter & Gamble,
Cincinnati OH 45202.**

PRINCIPAL DISPLAY PANEL - 354 ml Bottle Label

NEW *From the makers of* VICKS ® NyQuil®

ZzzQuil™

NIGHTTIME SLEEP-AID

Diphenhydramine HCl

- **Non-Habit Forming**
- **Warming Berry Flavor**

Not for treating Cold or Flu

See Warnings

Alcohol 10%

12 FL OZ (354 ml)

ZZZQUIL NIGHTTIME SLEEP-AID

diphenhydramine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37000-500
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**
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ALCOHOL (UNII: 3K9958V90M)	
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)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	

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:37000-500-06	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/08/2011	
2	NDC:37000-500-12	354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/08/2011	
3	NDC:37000-500-24	2 in 1 PACKAGE, COMBINATION	12/08/2011	
3	NDC:37000-500-12	354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:37000-500-36	3 in 1 PACKAGE, COMBINATION	12/08/2011	
4	NDC:37000-500-12	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 final	part338	12/08/2011	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 2/2021 The Procter & Gamble Manufacturing Company