

NIGHTTIME SLEEP AID- nighttime sleep aid liquid

Allegiant Health

446 - Nighttime Sleep Aid

Active ingredient(s)

Diphenhydramine HCl 50mg

In each 30mL dose cup or 2 tablespoons

Purpose

Nighttime sleep-aid

Use(s)

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

Warnings

Do not use

- Do not use n for children under 12 years of age.
- with any other product containing diphenhydramine, even one used on skin
- with any other drugs that cause drowsiness such as antihistamines and nighttime cough, cold/flu products

Ask a doctor before use if

- heart disease
- trouble urinating due to an enlarged prostate gland
- glaucoma
- a breathing problem such as asthma, emphysema or chronic bronchitis

Ask a doctor or pharmacist before use if

you are taking sedatives or tranquilizers or any other sleep aid

When using this product

- do not use more than directed
- avoid alcoholic beverages and other drugs that cause drowsiness
- drowsiness will occur
- be careful when driving a motor vehicle or operating machinery
- alcohol, sedatives, and tranquilizers may increase drowsiness

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 breastfeeding,

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

- take only as recommended
- use dosage cup or tablespoon

Age

Dose

Adults & children

12 years & over: One Dose = 2 tablespoons (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 23 mg
- dosage cup provided
- store at room temperature
- do not use if imprinted shrink band is missing or broken

Inactive ingredients

citric acid, FD&C Blue #1, FD&C Red #40, flavor, high fructose corn syrup, poloxamer 407, propylene glycol, purified water, sodium benzoate, sodium citrate, sodium saccharin

Questions/Comments

Call 1-888-952-0050 Monday through Friday 9AM to 5PM

Principal Display Panel

Drug Facts (continued)

Directions

- take only as recommended
- use dosage cup or tablespoon

| Age | Dose |
|-----------------------------------|---|
| Adults & children 12 years & over | One Dose = 2 tablespoons (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 23 mg
- dosage cup provided
- store at room temperature

Inactive ingredients
 citric acid, FD&C Blue #1, FD&C Red #40, flavor, high fructose corn syrup, poloxamer 407, propylene glycol, purified water, sodium benzoate, sodium citrate, sodium saccharin

Questions or comments?
 Call 1-888-952-0050 Monday through Friday 9AM - 5PM

*This product is not manufactured or distributed by Procter & Gamble, owner of the registered trademark ZzzQuil™

CH Manufactured for: **Allegiant Health**
 Deer Park, NY 11729

LB2119
 R0425
 LR-202 Rev 01

X003YSHVJJ
 HealthA2Z Nighttime ...Berry Flavored Liquid
 New

Lot:
 Exp:

HealthA2Z®

NDC 69168-446-39
 *Compare to active ingredient in ZzzQuil®

Nighttime Sleep Aid

Diphenhydramine HCl 50 mg

Relieves Occasional Sleeplessness



Berry Flavored Liquid
12 FL OZ (354 mL)

DO NOT USE IF IMPRINTED SHRINK BAND IS MISSING OR BROKEN

Drug Facts

| Active ingredient | Purpose |
|--|---------------------|
| (in each 30 mL dose cup or 2 tablespoons) Diphenhydramine HCl 50 mg | Nighttime sleep-aid |

Uses

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

Warnings

Do not use

- for children under 12 years of age.
- with any other product containing diphenhydramine, even one used on skin
- with any other drugs that cause drowsiness such as antihistamines and nighttime cough, cold/flu products

Ask a doctor before use if you have

- heart disease
- trouble urinating due to an enlarged prostate gland
- glaucoma
- a breathing problem such as asthma, emphysema or chronic bronchitis

Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers or any other sleep aid

When using this product

- do not use more than directed
- avoid alcoholic beverages and other drugs that cause drowsiness
- drowsiness will occur
- be careful when driving a motor vehicle or operating machinery
- alcohol, sedatives, and tranquilizers may increase drowsiness

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

Nighttime Sleep Aid

NIGHTTIME SLEEP AID

nighttime sleep aid liquid

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:69168-446 |
| 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) | |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | |
| FD&C RED NO. 40 (UNII: WZB9127XOA) | |
| HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S) | |
| POLOXAMER 407 (UNII: TUF2IVW3M2) | |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | |
| WATER (UNII: 059QF0KO0R) | |
| SODIUM BENZOATE (UNII: OJ245FE5EU) | |
| SODIUM CITRATE (UNII: 1Q73Q2JULR) | |
| SACCHARIN (UNII: FST467XS7D) | |

Product Characteristics

| | | | |
|-----------------|-------|---------------------|--|
| Color | | Score | |
| Shape | | Size | |
| Flavor | BERRY | Imprint Code | |
| Contains | | | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:69168-446-39 | 354 mL in 1 BOTTLE; Type 0: Not a Combination Product | 01/16/2024 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---------------------------|---|-----------------------------|---------------------------|
| OTC Monograph Drug | M010 | 01/16/2024 | |

