

SLEEP AID- diphenhydramine hcl solution
Wal-Mart Stores Inc

Equate 44-047

Active ingredient (in each 30 mL)

Diphenhydramine HCl 50 mg

Purpose

Nighttime sleep-aid

Uses

- for 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 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 right away.

Directions

- **do not take more than directed**
- mL = milliliter
- only use the dose cup provided
- take only one dose per day (24 hours)
- adults and children 12 years and over: 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 21 mg
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- use by expiration date on package

Inactive ingredients

anhydrous citric acid, FD&C blue #1, FD&C red #40, flavors, glycerin, high fructose corn syrup, propylene glycol, purified water, sodium benzoate, sodium chloride, sodium citrate dihydrate, sucralose, sucrose, xanthan gum

Questions or comments?

1-888-287-1915

Principal display panel

NDC 49035-704-02

Equate™

Compare to
VICKS® ZzzQuil®
Nighttime
Sleep-Aid
active
ingredient*

Nighttime Sleep-Aid

Diphenhydramine HCl
50 mg per 30 mL
Nighttime Sleep-Aid

- Non-habit forming
- Not for colds or
for pain
- Ages 12+

12 FL OZ (355 mL)

Cherry
Vanilla

F-047
ORG

Satisfaction guaranteed - Or we'll replace it or give you your money back. For questions or comments or to report an undesired reaction or side effect, please call **1-888-287-1915**.

DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716
PRODUCT OF CHINA

*This product is not manufactured or distributed by The Procter & Gamble Company, owner of the registered trademark VICKS® ZzzQuil® Nighttime Sleep-Aid.

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ORG052304702

B-047 ORG

**TAMPER EVIDENT: DO NOT USE IF PRINTED
NECK WRAP IS BROKEN OR MISSING**

NDX 49035-704-02



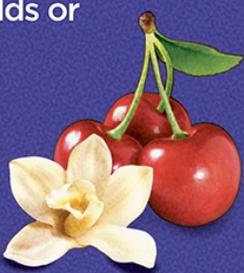
Compare to VICKS® ZzzQuil® Nighttime Sleep-Aid active ingredient*



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Cherry
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F-047
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PEEL BACK FOR COMPLETE DRUG FACTS

Drug Facts TAMPER EVIDENT: DO NOT USE IF PRINTED NECK WRAP IS BROKEN OR MISSING

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50844
ORG052304702
B-047 ORG

**NO PRINT / NO VARNISH AREA
LOT NO & EXP DATE**

Drug Facts (continued)

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Equate 44-047

SLEEP AID

diphenhydramine hcl solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-704
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)	
GLYCERIN (UNII: PDC6A3C0OX)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SUCROSE (UNII: C151H8M554)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	red (Maroon)	Score	
Shape		Size	
Flavor	CHERRY, VANILLA	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-704-02	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/18/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M010	03/18/2019	

Labeler - Wal-Mart Stores Inc (051957769)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	manufacture(49035-704) , pack(49035-704)

Revised: 3/2025

Wal-Mart Stores Inc