

SLEEP AID PLUS PAIN RELIEF- acetaminophen, diphenhydramine hcl solution
Walgreen Company

Walgreens 44-074 Sleep Aid Plus Pain Relief

Active ingredients (in each 30 mL)

Acetaminophen 1,000 mg
Diphenhydramine HCl 50 mg

Purpose

Pain reliever
Nighttime sleep-aid

Use

for the temporary relief of occasional minor aches and pains with accompanying sleeplessness

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- with any other product containing diphenhydramine, even one used on skin
- in children under 12 years of age
- if you have ever had an allergic reaction to this product or any of its ingredients

Ask a doctor before use if you have

- liver disease
- glaucoma

- a breathing problem such as emphysema or chronic bronchitis
- difficulty in urination due to enlargement of the prostate gland

Ask a doctor or pharmacist before use if you are

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers

When using this product

- drowsiness will occur
- avoid alcoholic beverages
- do not drive a motor vehicle or operate machinery

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.
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur

These could be signs of a serious condition.

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. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

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
- children under 12 years: do not use

Other information

- **each 30 mL contains:** sodium 12 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, propylene glycol,

purified water, sodium benzoate, sodium chloride, sodium citrate dihydrate, sorbitol, sucralose, xanthan gum

Questions or comments?

1-800-426-9391

Principal display panel

Walgreens

**Compare to the active ingredients
in Vicks® ZzzQuil® Night Pain††**

WALGREENS PHARMACIST RECOMMENDED†

Sleep Aid

+ Pain

Relief

ACETAMINOPHEN

PAIN RELIEVER

DIPHENHYDRAMINE HCl

NIGHTTIME SLEEP AID

Nighttime

- Non-habit forming
- Maximum strength pain reliever
- Ages 12 Years and Over

12 FL OZ (355 mL)

Midnight Berry
flavor

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

†Our pharmacists recommend the
Walgreens brand. We invite you to
compare to national brands.

††This product is not manufactured or
distributed by The Procter & Gamble
Company, owner of the registered
trademarks VICKS® ZzzQuil® NIGHT PAIN.

DISTRIBUTED BY: **WALGREEN CO.**
DEERFIELD, IL 60015

walgreens.com 1-800-925-4733

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NDC 0363-9074-02

50844 ORG052307402

Walgreens



Compare to the active ingredients in Vicks® ZzzQuil® Night Pain!

Sleep Aid + Pain Relief

ACETAMINOPHEN
PAIN RELIEVER
DIPHENHYDRAMINE HCl
NIGHTTIME SLEEP-AID

Nighttime

- Non-habit forming
- Maximum strength pain reliever
- Ages 12 Years and Over



12 FL OZ (355 mL)

Midnight Berry flavor

F-074 ORG REV-0424

PEEL BACK FOR COMPLETE DRUG FACTS

Drug Facts

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

Active ingredients (in each 30 mL)	Purpose
Acetaminophen 1,000 mg	Pain reliever
Diphenhydramine HCl 50 mg	Nighttime sleep-aid

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DEERFIELD, IL 60015
100% SATISFACTION GUARANTEED
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For more info

ITEM 692769 W00000-0000-0



NDC 0363-9074-02
W3ORG0324-F2
50844 ORG052307402
8-074 ORG

3 11917 04762 1

no print / no varnish area
lot no. & exp. date

Drug Facts (continued)

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- with any other product containing diphenhydramine, even one used on skin
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- liver disease
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Questions or comments?

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Walgreens 44-074

SLEEP AID PLUS PAIN RELIEF

acetaminophen, diphenhydramine hcl solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-9074
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	1000 mg in 30 mL
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)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	purple (Dark)	Score	
Shape		Size	
Flavor	BERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-9074-02	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/27/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	08/27/2022	

Labeler - Walgreen Company (008965063)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	manufacture(0363-9074) , pack(0363-9074)

Revised: 6/2025

Walgreen Company