VICKS ZZZQUIL NIGHT PAIN NIGHTTIME SLEEP-AID PAIN RELIEVER- diphenhydramine hcl, acetaminophen liquid
The Procter & Gamble Manufacturing Company

VICKS ® ZzzQuil™ NIGHT PAIN
NIGHTTIME SLEEP-AID
PAIN RELIEVER, Chill Mint Flavored

Drug Facts

Active ingredients (in each 30 mL)

Acetaminophen 1000 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 doses (30 mL each) in 24 hrs, which is the maximum daily amount for this product

• 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.
- 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

- liver disease
- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- difficulty in urination due to enlargement of the prostate gland
- a sodium-restricted diet

Ask a doctor or pharmacist before use if you are

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

When using this product

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

Stop use and ask a doctor if

- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- new symptoms occur
- sleeplessness persists continuously for more than 2 weeks. Insomnia may be a symptom of serious underlying medical illness.

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

Directions

• take only one dose (30 mL) per day (24 hours)
• only use the dose cup provided

adults & children 12 yrs & over	30 mL at bedtime
children under 12 yrs	do not use

Other information

- **each 30 mL dose cup contains:** sodium 88 mg
- store at no greater than 25°C and do not refrigerate

Inactive ingredients

alcohol, citric acid, D&C Yellow No. 10, FD&C Blue No. 1, flavor, polysorbate 20, propylene glycol, saccharin sodium, sodium benzoate, sodium chloride, sodium citrate, sorbitol, sucralose, water, xanthan gum

Questions?

1-877-881-5813

TAMPER EVIDENT: DO NOT USE IF PRINTED SHRINKBAND IS BROKEN OR MISSING.

DIST. BY PROCTER & GAMBLE,

CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 354 ml bottle

VICKS ®

ZzzQuil™
NIGHT PAIN

NIGHTTIME SLEEP-AID

PAIN RELIEVER

Diphenhydramine HCl

Acetaminophen

Fall Asleep Fast

Max Strength Pain Reliever

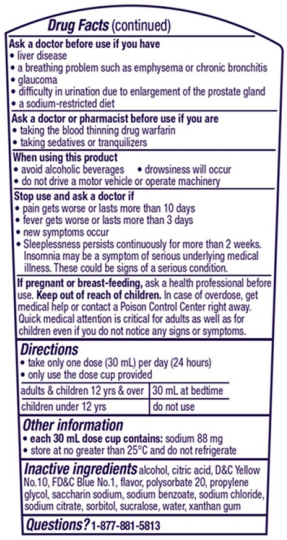
Non-Habit Forming

Chill Mint Flavored

Not for colds

Alcohol 10%

12 FL OZ (354 ml)



VICKS ZZZQUIL NIGHT PAIN NIGHTTIME SLEEP-AID PAIN RELIEVER

diphenhydramine hcl, acetaminophen liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84126-372
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
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	1000 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SORBITOL (UNII: 506T60A25R)	

SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics			
Color	blue	**Score**	
Shape		**Size**	
Flavor	MINT	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84126-372-12	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/03/2026	
2	NDC:84126-372-24	2 in 1 CARTON	02/03/2026	
2		354 mL in 1 BOTTLE; 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	02/03/2026	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 1/2026 The Procter & Gamble Manufacturing Company