

**VICKS PAINQUIL PM PAIN RELIEVER PLUS NIGHTTIME SLEEP-AID-
acetaminophen, diphenhydramine hcl liquid
The Procter & Gamble Manufacturing Company**

Vicks® PainQuil™ PM PAIN RELIEVER + NIGHTTIME SLEEP-AID

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 3 doses (30 mL each) in 24 hours, 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

Sore throat warning: If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

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

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
- redness or swelling is present
- 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.

When using this product

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

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
- only use as directed

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

Other information

- **each 30 mL contains:** sodium 89 mg
- do not exceed 25°C and do not refrigerate.

Inactive ingredients

alcohol, anhydrous citric acid, FD&C Blue No. 1, FD&C Red No. 40, 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®

PainQuil™ PM

PAIN RELIEVER + NIGHTTIME SLEEP-AID

Acetaminophen

Diphenhydramine HCl

STARTS WORKING FAST

FOR RELIEF OF:

ACHES | PAINS

HEADACHE | SORE THROAT

+ OCCASSIONAL SLEEPLESSNESS

MIDNIGHT CHERRY FLAVORED

Alcohol 10%

12 FL OZ (354 ml)



Drug Facts (continued)

Directions

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

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

Other information

- each 30 mL dose contains: sodium 89 mg
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Inactive ingredients alcohol, anhydrous citric acid, FD&C Blue No. 1, FD&C Red No. 40, flavor, polysorbate 20, propylene glycol, saccharin sodium, sodium benzoate, sodium chloride, sodium citrate, sorbitol, sucralose, water, xanthan gum

QUESTIONS? 1-877-881-5813

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.
- 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

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
- redness or swelling is present
- 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.

When using this product

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

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.



VICKS PAINQUIL PM PAIN RELIEVER PLUS NIGHTTIME SLEEP-AID

acetaminophen, diphenhydramine hcl liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69423-834
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
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALCOHOL (UNII: 3K9958V90M)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
WATER (UNII: 059QF0KO0R)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	

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

Product Characteristics			
Color	purple	Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-834-12	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/07/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M010	05/07/2024	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 4/2024

The Procter & Gamble Manufacturing Company