

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

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**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

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

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**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)**

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**Drug Facts (continued)**

**Do not use**

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- for children under 12 years of age
- with any other product containing diphenhydramine, even one used on skin

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- 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**

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**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

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69423-834
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	50 mg in 30 mL
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	1000 mg in 30 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>POLYSORBATE 20</b> (UNII: 7T1F30V5YH)	
<b>SACCHARIN SODIUM</b> (UNII: SB8ZUX40TY)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>SUCRALOSE</b> (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	M013	05/07/2024	

**Labeler** - The Procter & Gamble Manufacturing Company (004238200)

Revised: 12/2025

The Procter & Gamble Manufacturing Company