VICKS PAINQUIL PM PAIN RELIEVER PLUS NIGHTTIME SLEEP-AIDacetaminophen, 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 HCI 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 &	30 mL at
over	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

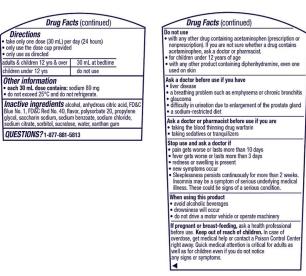
+ OCCASSIONAL SLEEPLESSNESS

MIDNIGHT CHERRY FLAVORED

Alcohol 10%

12 FL OZ (354 ml)







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: 36209ITL9D) (ACETAMINOPHEN - UNII: 36209ITL9D)	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	M013	05/07/2024	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 12/2024 The Procter & Gamble Manufacturing Company