

**VICKS PAINQUIL AND VICKS PAINQUIL PM PAIN RELIEVER- acetaminophen,  
diphenhydramine hcl**

**The Procter & Gamble Manufacturing Company**

-----  
**Vicks® PainQuil™ / Vicks® PainQuil™ PM  
PAIN RELIEVER Convenience Pack**

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

**Uses**

- 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**

Vicks ® PainQuil™ PAIN RELIEVER

### ***Drug Facts***

#### **Active ingredients (in each 30 mL)**

Acetaminophen 1000 mg

#### ***Purpose***

Pain reliever

#### **Uses**

for the temporary relief of minor aches and pains associated with:

- sore throat
- headache
- muscular aches
- backache

#### **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.

### **Ask a doctor before use if you have**

- liver disease
- a sodium-restricted diet

### **Ask a doctor or pharmacist before use if you are**

- taking the blood thinning drug warfarin

### **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.

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 as directed
- do not exceed 3 doses (30 mL each) per day (24 hours)
- only use the dose cup provided

---

adults & children 12 yrs & over  
children under 12 yrs

---

30 mL every 6 hours  
do not use

---

### **Other information**

- **each 30 mL contains:** sodium 93 mg
- do not exceed 25°C and do not refrigerate.

### **Inactive ingredients**

alcohol, anhydrous citric acid, 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 - Convenience Pack**

Vicks<sup>®</sup> PainQuil<sup>™</sup> PAIN RELIEVER / Vicks<sup>®</sup> PainQuil<sup>™</sup> PM PAIN RELIEVER + NIGHTTIME SLEEP-AID  
MAX STRENGTH‡

VALUE PACK

‡Maximum strength dose of active ingredients per dosing interval, only use as directed.

Vicks<sup>®</sup> PainQuil<sup>™</sup> PAIN RELIEVER

STARTS WORKING FAST FOR RELIEF OF:

ACHES

PAINS

HEADACHE

SORE THROAT

BLACK CHERRY FLAVORED

Alcohol 10%

Vicks<sup>®</sup> PainQuil<sup>™</sup> PM PAIN RELIEVER + NIGHTTIME SLEEP-AID

STARTS WORKING FAST FOR RELIEF OF:

ACHES  
 PAINS  
 HEADACHE  
 SORE THROAT  
 +  
 OCCASSIONAL SLEEPLESSNESS  
 MIDNIGHT CHERRY FLAVORED  
 Alcohol 10%

2 BOTTLES - 1 PAINQUIL/1 PAINQUIL PM 12 FL OZ (354 mL) EACH; TOTAL 24 FL OZ (708 mL)



## VICKS PAINQUIL AND VICKS PAINQUIL PM PAIN RELIEVER

acetaminophen, diphenhydramine hcl kit

### Product Information

**Product Type**

HUMAN OTC DRUG

**Item Code (Source)**

NDC:69423-849

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-849-24	1 in 1 PACKAGE; Type 0: Not a Combination Product	05/07/2024	
2	NDC:69423-849-36	1 in 1 PACKAGE; Type 0: Not a Combination Product	03/17/2025	

## Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	354 mL
Part 2	1 BOTTLE	354 mL
Part 3	1 BOTTLE	354 mL in 2

## Part 1 of 3

### VICKS PAINQUIL PAIN RELIEVER

acetaminophen liquid

## Product Information

Item Code (Source)	NDC:69423-833
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

## Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
WATER (UNII: 059QF0KO0R)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SORBITOL (UNII: 506T60A25R)	

<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	

**Product Characteristics**

<b>Color</b>	red	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	CHERRY	<b>Imprint Code</b>	
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-833-12	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	M013	05/07/2024	

**Part 2 of 3**

**VICKS PAINQUIL PM PAIN RELIEVER PLUS NIGHTTIME SLEEP-AID**  
 acetaminophen, diphenhydramine hcl liquid

**Product Information**

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

**Inactive Ingredients**

Ingredient Name	Strength
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<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>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>TRISODIUM CITRATE DIHYDRATE</b> (UNII: B22547B95K)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	

### Product Characteristics

<b>Color</b>	purple	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	CHERRY	<b>Imprint Code</b>	
<b>Contains</b>			

### 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		
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	M013	05/07/2024	

### Part 3 of 3

## VICKS PAINQUIL PM PAIN RELIEVER PLUS NIGHTTIME SLEEP-AID

acetaminophen, diphenhydramine hcl liquid

### Product Information

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

### Inactive Ingredients

Ingredient Name	Strength
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<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>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>TRISODIUM CITRATE DIHYDRATE</b> (UNII: B22547B95K)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	

### Product Characteristics

<b>Color</b>	purple	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	CHERRY	<b>Imprint Code</b>	
<b>Contains</b>			

### 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		
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	M013	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