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	30 mL every 6 hours
children under 12 yrs	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?

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

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CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - Convenience Pack

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

VALUE PACK

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

Vicks ®PainQuil™ PAIN RELIEVER

STARTS WORKING FAST FOR RELIEF OF:

ACHES

PAINS

HEADACHE

SORE THROAT

BLACK CHERRY FLAVORED

Alcohol 10%

Vicks ®PainQuil™ 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)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics			
Color	red	**Score**	
Shape		**Size**	
Flavor	CHERRY	**Imprint Code**	
Contains			

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	
Item Code (Source)	NDC:69423-834
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
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	50 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
WATER (UNII: 059QF0KO0R)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
XANTHAN GUM (UNII: TTV12P4NEE)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALCOHOL (UNII: 3K9958V90M)	

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		
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

Item Code (Source)	NDC:69423-834
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
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	50 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
WATER (UNII: 059QF0KO0R)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
XANTHAN GUM (UNII: TTV12P4NEE)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALCOHOL (UNII: 3K9958V90M)	

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		
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: 2/2025 The Procter & Gamble Manufacturing Company