

**VICKS DAYQUIL HIGH BLOOD PRESSURE COLD AND FLU- acetaminophen,
dextromethorphan liquid**
The Procter & Gamble Manufacturing Company

VICKS® DayQuil™
HIGH BLOOD PRESSURE COLD & FLU

Drug Facts

Active ingredients (in each 30 mL)

Acetaminophen 650 mg
Dextromethorphan HBr 20 mg

Purpose

Pain reliever/fever reducer
Cough suppressant

Uses

temporarily relieves common cold/flu symptoms:

- cough due to minor throat and bronchial irritation
- sore throat
- headache
- minor aches and pains
- fever

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take

- more than 4 doses (30 mL each dose) in 24 hrs, 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.
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

Ask a doctor before use if you have

- liver disease
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema

Ask a doctor or pharmacist before use if you are

- taking the blood thinning drug warfarin

Stop use and ask a doctor if

- pain or cough gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- cough comes back or occurs with rash or headache that lasts.

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
- only use dose cup provided
- do not exceed 4 doses per 24 hrs

adults & children 12 yrs & over
 children 4 to under 12 yrs
 children under 4 yrs

30 mL per dose every 4 hrs
 ask a doctor
 do not use

Other information

- **each 30 mL contains:** sodium 29 mg
- store at no greater than 25°C and do not refrigerate

Inactive ingredients

citric acid, FD&C yellow no. 6, flavor, glycerin, propylene glycol, saccharin sodium, sodium benzoate, sodium citrate, sorbitol, sucralose, water, xanthan gum

Questions?

1-800-362-1683

TAMPER EVIDENT:

Do not use if printed shrinkband seal around the neck is broken or missing

DIST. BY

PROCTER & GAMBLE

CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 236 ml Bottle Label

VICKS®

DayQuil™

HIGH BLOOD PRESSURE

COLD & FLU

Acetaminophen, Dextromethorphan HBr

Headache, Fever, Sore Throat, Minor Aches & Pains

Cough

SUGAR & ALCOHOL FREE

DECONGESTANT FREE

Non-Drowsy

8 FL OZ (236 mL)



Drug Facts (continued)

Stop use and ask a doctor if

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- cough comes back, or occurs with rash or headache that lasts.

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← **PEEL BACK FOR DRUG FACTS**
ALSO SOLD AS DAYQUIL FOR PEOPLE WITH DIABETES

Drug Facts

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Patents: www.pg.com/patents

PARENTS: Learn about teen medicine abuse
www.StopMedicineAbuse.org

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VICKS DAYQUIL HIGH BLOOD PRESSURE COLD AND FLU

acetaminophen, dextromethorphan liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69423-972
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg in 30 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL (UNII: 506T60A25R)	

XANTHAN GUM (UNII: TTV12P4NEE)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

Product Characteristics

Color	orange	Score	
Shape		Size	
Flavor	CITRUS	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-972-06	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/12/2022	
2	NDC:69423-972-08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/12/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/12/2022	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 12/2025

The Procter & Gamble Manufacturing Company