

COLD AND FLU RELIEF- acetaminophen dextromethorphan hbr phenylephrine hcl liquid

QUALITY CHOICE (Chain Drug Marketing Association)

Drug Facts

Active ingredients (in each 15 mL)

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Phenylephrine HCl 5 mg

Purpose

Pain reliever/fever reducer

Cough suppressant

Nasal decongestant

Uses

- temporarily relieves common cold and flu symptoms:
 - minor aches and pains
 - headache
 - sore throat
 - nasal congestion
 - fever
 - sinus pressure
 - cough due to minor throat and bronchial irritation

Warnings

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

- adult take more than 4 doses (30 mL each) of acetaminophen in 24 hours, which is the maximum daily amount
- child takes more than 4 doses (15 mL each) in 24 hours
- taken with other drugs containing acetaminophen
- adult has 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 is 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
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough as occurs with smoking, asthma, or emphysema
- cough that occurs with too much phlegm (mucus)

Ask a doctor or pharmacist before use if user is

taking the blood thinning drug warfarin

When using this product

do not exceed recommended dosage.

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- pain, nasal congestion, or cough gets worse or lasts more than 5 days (children) or 7 days (adults)
- 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.

Overdose warning: Taking more than the recommended dose (overdose) can cause serious liver damage. In case of overdose, get medical help or contact a Poison Control

Center (1-800-222-1222) right away. Quick medical attention is critical for adults and for children even if you do not notice any signs or symptoms

Directions

- **do not take more than directed (see Overdose warning)**
- do not take more than 4 doses in any 24-hour period
- measure only with dosing cup provided. Do not use any other dosing device
- mL = milliliter
- keep dosing cup with product

adults and children 12 years and over	30 mL every 4 hours
children 6 to under 12 years	15 mL every 4 hours
children 4 to under 6 years	ask a doctor
children under 4 years	do not use

- **when using other Daytime or Nighttime products, carefully read each label to ensure correct dosing**

Other information

- each 15 mL contains: sodium 12 mg
- store between 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

citric acid, FD&C Yellow #6, flavor, glycerin, propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium chloride, sodium citrate, sorbitol sucralose, xanthan gum

Questions or comments?

Call 1-800-935-2362 Monday-Friday 9AM-5PM EST

Principal Display Panel

*Compare to the active ingredients in Vicks® DayQuil™ Cold & Flu

Multi-Symptom

Daytime

Cold & Flu Relief

Acetaminophen

Pain reliever/Fever reducer

Dextromethorphan HBr

Cough suppressant

Phenylephrine HCl

Nasal decongestant

Relieves

Aches, fever & sore throat

Cough

Nasal congestion

Sinus pressure

For ages 6 years and over

Alcohol-free

Antihistamine-free

FL OZ (mL)

*This product is not manufactured or distributed by The Procter & Gamble Company. Vicks® and DayQuil™ are registered trademarks of The Procter & Gamble Company.

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL AROUND DOSAGE CUP OR UNDER CAP IS BROKEN OR MISSING.

DISTRIBUTED BY:

Product Label



NDC 83324-168-08

*Compare to the Active Ingredients in Vicks® DayQuil™ Cold & Flu

Multi-Symptom Daytime

Cold & Flu Relief

Acetaminophen

Pain Reliever/Fever Reducer
Dextromethorphan HBr
Cough Suppressant
Phenylephrine HCl
Nasal Decongestant



Relieves:

Aches, fever & sore throat
Cough
Nasal congestion
Sinus pressure

For Ages 6 Years & Over
Alcohol Free
Antihistamine Free

Distributed by CDMA, Inc.
Novi, MI 48375
www.qualitychoice.com
Questions: 800-935-2362



8 fl oz (237 mL)

Drug Facts

Active ingredients (in each 15 mL) Purposes

Acetaminophen 325 mg.....Pain reliever/fever reducer
Dextromethorphan HBr 10 mg.....Cough suppressant
Phenylephrine HCl 5 mg.....Nasal decongestant

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PARENTS:

Learn about teen medicine abuse
www.StopMedicineAbuse.org

(Optional precautionary statement for Dextromethorphan HBr. Specific to customer's labeling guide)

Distributed by: (Specific to customer's labeling guide. Must state either: manufacturer, packer, or distributor.)

PLD-B380E LB009716



PEEL CORNER FOR MORE DRUG FACTS

Drug Facts (continued)

Uses

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PEEL CORNER FOR MORE DRUG FACTS

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Ask a doctor before use if you have

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema
- cough that occurs with too much phlegm (mucus)

Ask a doctor or pharmacist before use if you are taking the blood thinning drug

Drug Facts (continued)

When using this product, do not exceed recommended dosage.

Stop use and ask a doctor if

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- pain, nasal congestion, or cough gets worse, or lasts more than 5 days (children) or 7 days (adults)
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Drug Facts (continued)

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Inactive ingredients citric acid, FD&C yellow #6, flavor, glycerin, propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium chloride, sodium citrate, sorbitol, sucralose, xanthan gum

Questions or comments?

Call 1-800-XXX-XXXX DAYS & TIMES & ZONE

you are taking the blood thinning drug
warfarin.

any signs or symptoms.

QUALITY CHOICE Multi-Symptom Daytime Cold & Flu Relief

COLD AND FLU RELIEF

acetaminophen dextromethorphan hbr phenylephrine hcl liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83324-168
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg in 15 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 15 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-168-08	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/31/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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OTC Monograph Drug M012

07/31/2024

Labeler - QUALITY CHOICE (Chain Drug Marketing Association) (011920774)

Revised: 11/2025

QUALITY CHOICE (Chain Drug Marketing Association)