COLD AND FLU RELIEF- acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride liquid Gobrands, Inc. (Goodnow)

Drug Facts

Active ingredients (in each 15 mL)

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Phenylephrine HCI 5 mg

Purpose

Pain reliever/fever reducer

Cough suppressant

Nasal decongestant

Uses

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

Warnings

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

- 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
- blistere
- 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
- a sodium-restricted diet
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough as occurs with smoking, asthma, or emphysema

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)
- new symptoms occur
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- 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 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 24-hours period
- measure only with dosing cup provided. Do not use any other dosing device.
- mL = milliliter
- keep dosing cup with product

adults and children12 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 **1-877-753-3935** Monday-Friday 9AM-5PM EST

Principal Display Panel

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

NON-DROWSY

Cold & Flu

Day Relief

Acetaminophen

Dextromethorphan HBr

Phenylephrine HCI

Pain Reliever

Fever Reducer

Cough Suppressant

Nasal Decongestant

Alcohol-free / Anthihistamine-free

Multi-Symptom Relief

Headache & Fever

Sore Throat

Minor Aches & Pains

Nasal Congestion

Cough

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.

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PHILADELPHIA, PA 19123

Product Label



COLD AND FLU RELIEF

acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride liquid

Product Information	roduct Information		
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82501-4661
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	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)		

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	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:82501- 4661-2	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/29/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	04/29/2022	

Labeler - Gobrands, Inc. (Goodnow) (057499049)

Revised: 4/2024 Gobrands, Inc. (Goodnow)