COLD AND FLU RELIEF- acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride liquid Walgreens

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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
 - 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)
- 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 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 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 & Flutt

DAYTIME • NON-DROWSY

Cold & Flu

ACETAMINOPHEN 325 mg / PAIN RELIEVER / FEVER REDUCER

DEXTROMETHORPHAN HBr 10 mg / COUGH SUPPRESSANT

PHENYLEPHRINE HCI 5 mg / NASAL CONGESTION

Multi-Symptom

- Relieves aches, fever & sore throat , cough & nasal congestant
- For ages 6 years & 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: WALGREEN CO.

200 WILMOT RD., DEERFIELD, IL 60015

Product Label

Drug Facts

Dextromethorphan HBr 10 mg Phenylephrine HCl 5 mg	
Uses temporarily relieves common co minor aches and pains headache nasal congestion fever cough due to minor throat and bronchial in	sore throat
Warnings Liver warning: This product contains acetan damage may occur if: a dult takes more each) of acetaminophen in 24 hours, which is amount b child takes more than 4 doses (hours taken with other drugs containing adult has 3 or more alcoholic drinks every product Allergy alert: Acetaminophen may cause see Symptoms may include: skin reddening If a skin reaction occurs, stop use and seek n Sore throat warning: If sore throat is severe 2 days, is accompanied or followed by fever, nausea, or vomiting, consult a doctor prompti	than 4 doses (30 mL s the maximum daily 15 mL each) in 24 ng acetaminophen day while using this vere skin reactions. ■ blisters ■ rash nedical help right away. , persists for more than headache, rash,
Do not use with any other drug containi (prescription or nonprescription). If you are no contains acetaminophen, ask a doctor or pha	ot sure whether a drug

Tompare to the active ingredients in Vicks® DayQuil® Cold & Flu"

NDC 0363-9974-08

Cold & Flu

ACETAMINOPHEN 325 mg / PAIN RELIEVER /

FEVER REDUCER DEXTROMETHORPHAN HBr 10 mg / COUGH SUPPRESSANT PHENYLEPHRINE HCI 5 mg / NASAL DECONGESTANT

Multi-Symptom

- Relieves aches, fever, sore throat, cough & nasal congestion
- For ages 6 years & over
- Antihistamine free
- Alcohol free

8 FL 0Z (237 mL)

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL Around Dosage cup or under cap is broken or missing



Our pharmacists recommend the Walgreens brand. We invite you to compare to

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Drug Facts (continued)	Drug Facts (continued)		
If you are now taking a prescription monoamine addase inhibitor (MAD) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stapping the MADI drug. If you do not know if your prescription drug contains an MADI, ask a doctor or pharmacist	dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Cubic medical attention is critical for adults as well as 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 m measure only with dosing cup provided. Do not use any other dosing device. New Cosing cup with product m inter mainter		
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cough such as occurs with smoking, asthma, or emphysema in cough that occurs with too much philegm (mucus)	adults and children 12 years and over	30 mL every 4 hours	
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Keep out of reach of children. Overdose warning: Taking more than the recommended	Questions or comments? Call 1-877-753-3535 Monday-Friday 94	N-5PM EST	

WALGREENS Daytime Cold & Flu

COLD AND FLU RELIEF

acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-9974
Route of Administration	ORAL		
Active Ingredient/Active	Majaty		
Active Ingredient/Active Moiety			

	Ing	redient Name		Basis of Stre	ngth	Strength
ACETAMINOPHE	N (UNII: 362C	9ITL9D) (ACETAMINOPHEN - UNII:36	209ITL9D)	ACETAMINOPHEN		325 mg in 15 mL
DEXTROMETHO (DEXTROMETHOR		OBROMIDE (UNII: 9D2RTI9KYH) 855X3ROTS)		DEXTROMETHORPHAN HYDROBROMIDE		10 mg in 15 mL
PHENYLEPHRIN UNII:1WS297W6M		ORIDE (UNII: 04JA59TNSJ) (PHENYL	EPHRINE -	PHENYLEPHRINE HYDROCHLORIDE		5 mg in 15 mL
Inactive Ing	redients					
		Ingredient Name			Strength	
FD&C YELLOW	NO.6 (UNII: H	177VE193A8)				
GLYCERIN (UNII:	PDC6A3C0OX)				
PROPYLENE GL	(COL (UNII: 6	DC9Q167V3)				
WATER (UNII: 05	9QF0KO0R)					
SORBITOL (UNII:	506T60A25R)					
SACCHARIN SO	-					
SODIUM BENZO		•				
SODIUM CHLOR						
SODIUM CITRAT						
XANTHAN GUM						
SUCRALOSE (UN						
ANHYDROUS CI	RIC ACID (U	NII: XF417D3PSL)				
Packaging						
# Item Code		Package Description	M	larketing Start Date	Mark	eting End Date
1 NDC:0363- 9974-08	237 mL in 1 Combination	BOTTLE, PLASTIC; Type 0: Not a Product	09	/30/2022		
Marketing	lnform	ation				
	Ann	lication Number or Monogra	oh Ma	arketing Start	Mark	eting End
Marketing Category	Ahb	Citation		Date		Date

Labeler - Walgreens (008965063)

Revised: 4/2023

Walgreens