WALGREEN DAYTIME COLD AND FLU RELIEF- acetaminophen, dextromethorphan hydrobromide, guaifenesin and phenylephrine hcl liquid WALGREENS CO.

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.

Walgreen DayQuil Severe Cold & Flu Honey

Drug Facts

Active ingredients (in each 15 mL)

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Guaifensin 200 mg

Phenylephrine HCl 5 mg

Purpose

Pain reliever/fever reducer Cough suppressant

Expectorant

Nasal decongestant

Uses

temporarily relieves common cold/flu symptoms:

- nasal congestion
- sinus congestion & pressure
- cough due to minor throat & bronchial irritation
- minor aches & pains
- headache
- fever
- sore throat
- reduces swelling of nasal passages
- temporarily restores freer breathing through the nose
- promotes nasal and/or sinus drainage
- helps loosen phlegm (mucus) and thin bronchial secretions torid the bronchial passageways of bothersome mucus and make coughs more productive.

Warnings

Liver warning

This product contains acetaminophen. Severe liver damage may occur if

- adult takes more than 4 doses (30 mL each) in 24 hrs, which is the maximum daily amount for this product
- child takes more than 4 doses (15 mL each) in 24 hrs, which is the maximum daily amount for this product
- 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 enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- persists or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema

Ask a doctor or pharmacist before use if you are

taking the blood thinning drug warfarin

When using this product,

do not use more than directed

Stop use and ask a doctor if

- you get nervous, dizzy or sleepless
- pain, nasal congestion or cough gets worse or lasts more than 5 days (children) 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.

In case of overdose, get medical help or contact a Poison Control Center right away at 1-800-222-1222. 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 the dose cup provided
- do not exceed 4 doses per 24 hrs

adults & children 12 yrs & over 30 mL every 4 hrs

children 6 to under 12 yrs 15 mL every 4 hrs children 4 to under 6 yrs ask a doctor children under 4 yrs do not use

Other information

- each 15 mL contains: sodium 9 mg
- Store at room temperature
- Do not refrigerate.

Inactive ingredients

citric acid, D&C Yellow No. 10, edetate disodium, FD&C Green No. 3, FD&C Red No. 40, flavors, glycerin, menthol, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose, xanthan gum

Questions?

1-866-467-2748

*This product is not manufactured or distributed by Procter & Gamble, distributer of

Vicks[®] Dayquil[™] Severe Cold & Flu Honey Flavour

Distributed. by:

PRINCIPAL DISPLAY PANEL - 354 mL Bottle Label

*Compare to the active ingredients in Vicks® Dayquil[™] Severe Cold & Flu Honey Flavour NDC 0363-6110-12

DayTime

Cold & Flu

Relief

Acetaminophen - Pain reliever/Fever reducer

Guaifensin - Expectorant

Phenylephrine HCl - Nasal Congestion

Dextromethorphan HBr – Cough suppressant

- Headache, Fever, Sore Throat, Minor Aches & Pains •
- Chest Congestion, •
- Thins & Loosens Mucus
- Nasal Congestion, Sinus Pressure
- Cough •

HONEY FLAVOR

Naturally and Artificially Flavored

12 FL OZ (354 ml)



*Compare to the active ingredients in Vicks[®] Dayquil[™] Severe Cold & Flu Honey Flavor NDC 0363-6110-12

DayTime Cold & Flu

Acetaminophen - Pain reliever/Fever reducer Guaifenesin - Expectorant Phenylephrine HCI - Nasal decongestant Dextromethorphan HBr - Cough suppressant

- · Headache, Fever, Sore Throat,

- · Nasal Congestion, Sinus Pressure

Honey Flavor 12 FL OZ (354 ml)

Active ingredient	s (in each 15 mL)	Purpos
Dextromethorphan HBr Guaifenesin 200 mg	10 mg	Cough suppressar Expectoran
congestion & pressure & pains • headache • fev • temporarily restores fr drainage • helps loosen	eves common cold/flu symptoms: • cough due to minor throat & bronc er • sore throat • reduces swelling of eer breathing through the nose • pro philegm (mucus) and thin bronchilal of bothersome mucus and make co	chial irritation • minor aches of nasal passages omotes nasal and/or sinus secretions to rid the
adult takes more than amount for this product the maximum daily amo- taken with other drugs every day while using th Allergy Alert: Acetamin Skin reddening - Bliste If a skin reaction occurs Sore throat warning: If s or followed by fever, hea	containing acetaminophen • adult h is product priner may cause severe skin reaction rs • Rash stop use and seek medical help rig ore throat is severe, persists for more dache, rash, nausea, or vomitting, c	ich is the maximum daily mL each) in 24 hrs, which i las 3 or more alcoholic drink ons. Symptoms may include ht away e than 2 days, is accompaniec onsult a doctor promptly.
If you are not sure whet If you are now taking a drugs for depression, po 2 weeks after stopping to 2 weeks after stopping to 3 weeks after stopping to 3 weeks after stopping to 3 weeks after stopping to 3 weeks after stopping to 4 weeks after stopping to	er drug containing acetaminophen (p her a drug contains acetaminophen, prescription monoamine oxidase ir sychiatric or emotional conditions, o he MAOI drug. If you do not know if doctor or pharmacist before taking	, ask a doctor or pharmacist. hibitor (MAOI) (certain or Parkinson's disease), or fo f your prescription drug
Ask a doctor before use • thyroid disease • diabe that occurs with too mu	If you have • liver disease • heart of tes • trouble urinating due to enlarg of phlegm (mucus) • persistent or of hronic bronchitis, or emphysema	disease • high blood pressu ed prostate gland • cough

WALGREEN DAYTIME COLD AND FLU RELIEF

acetaminophen, dextromethorphan hydrobromide, guaifenesin and phenylephrine hcl liquid

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:036	3-6110
Route of Administration	ORAL				
Active Ingredient/Active	Moiety				
Ingre	dient Name		Basis of Stre	ength	Strengt
ACETAMINOPHEN (UNII: 36209IT	L9D) (ACETAMINOPHEN - UN	III:362O9ITL9D)	ACETAMINOPHEN		325 mg in 15 mL
DEXTROMETHORPHAN HYDROB (DEXTROMETHORPHAN - UNII:7355		1)	DEXTROMETHORPH HYDROBROMIDE	IAN	10 mg in 15 mL
PHENYLEPHRINE HYDROCHLOR UNII:1WS297W6MV)	IDE (UNII: 04JA59TNSJ) (PHE	ENYLEPHRINE -	PHENYLEPHRINE HYDROCHLORIDE		5 mg in 15 mL
GUAIFENESIN (UNII: 495W7451VC) (GUAIFENESIN - UNII:495V	/7451VQ)	GUAIFENESIN		200 mg in 15 mL
Inactive Ingredients					
	Ingredient Name			St	trength
CITRIC ACID MONOHYDRATE (U					
D&C YELLOW NO. 10 (UNII: 355					
EDETATE DISODIUM (UNII: 7FLDS					
FD&C GREEN NO. 3 (UNII: 3P3ON					
FD&C RED NO. 40 (UNII: WZ B912	27XOA)				
GLYCERIN (UNII: PDC6A3C0OX)					
MENTHOL, UNSPECIFIED FORM					
PROPYLENE GLYCOL (UNII: 6DC9	Q167V3)				
WATER (UNII: 059QF0KO0R)					
SODIUM BENZOATE (UNII: OJ245					
SODIUM CITRATE, UNSPECIFIE	D FORM (UNII: 1Q73Q2JULF	()			
SORBITOL (UNII: 506T60A25R)					
SUCRALOSE (UNII: 96K6UQ3ZD4)					
XANTHAN GUM (UNII: TTV12P4NE	E)				
Product Characteristics					
Color	brown Sc	ore			
Shape		ze			
Flavor		print Code			
Contains					
Contains					
Packaging					
i ackaging					

#	Item Code	Package Description	Marketing Start Date	Date
1		354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2021	
Μ	larketing	Information		
Μ	larketing Marketing Category	Information Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
	Marketing	Application Number or Monograph Citation	-	-

Labeler - WALGREENS CO. (008965063)

Revised: 2/2021

WALGREENS CO.