VICKS DAYQUIL KIDS COLD COUGH PLUS MUCUS- guaifenesin, phenylephrine hydrochloride and dextromethorphan hydrobromide liquid The Procter & Gamble Manufacturing Company

Vicks DayQuil Kids Honey Cold & Cough + Mucus Drug Facts

Active ingredients (in each 15 mL)

Dextromethorphan HBr 10 mg
Guaifenesin 100 mg
Phenylephrine HCl 5 mg

Purpose

Cough suppressant

Expectorant

Nasal decongestant

Uses

temporarily relieves common cold symptoms:

- nasal congestion
- sinus congestion & pressure
- cough due to minor throat & bronchial irritation
- cough to help you sleep
- 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 to rid the bronchial passageways of bothersome mucus and make coughs more productive..

Warnings

Do not use

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

- heart disease
- high blood pressure
- thyroid disease

- diabetes
- trouble urinating due to enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis or emphysema

When using this product, do not use more than directed.

Stop use and ask a doctor if

- you get nervous, dizzy or sleepless
- symptoms do not improve within 7 days or occur with a fever
- cough persists for more than 7 days, comes back or occurs with a fever, rash or persistent headache

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.

Directions

- take only as directed
- only use the dose cup provided
- do not exceed 4 doses per 24 hrs

adults & children 12 yrs &	30 mL every 4
over	hrs
children 6 to under 12 yrs	15 mL every 4
crimar err o to arraer 12 yrs	hrs
children 4 to under 6 yrs	ask a doctor
children under 4 yrs	do not use

Other information

- each 15 mL contains: sodium 14 mg
- store at no greater than 25°C.

Inactive ingredients

citric acid, D&C Yellow No. 10, FD&C Green No. 3, FD&C Red No. 40, FD&C Yellow No. 6, flavor (with honey), 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™

Kids

HONEY

COLD & COUGH + MUCUS

Guaifenesin, Phenylephrine HCl, Dextromethorphan HBr

Chest Congestion Nasal Congestion Cough

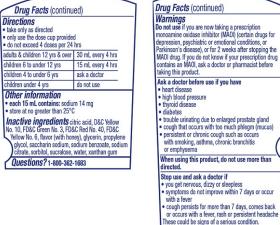
AGES 6+

8 FL OZ (236 mL)

Non-Drowsy

FREE OF ALCOHOL & ACETAMINOPHEN





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professional before use. **Keep out of reach of**children. In case of overdose, get medical help or
contact a Poison Control Center right away.



VICKS DAYQUIL KIDS COLD COUGH PLUS MUCUS

guaifenesin, phenylephrine hydrochloride and dextromethorphan hydrobromide liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69423-946	
Route of Administration	ORAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
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	
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 15 mL	

Inactive Ingredients	
Ingredient Name	Strength
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
XANTHAN GUM (UNII: TTV12P4NEE)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	

Product Characteristics			
Color	brown	Score	
Shape		Size	
Flavor	HONEY	Imprint Code	
Contains			

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:69423- 946-08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/08/2021	

Marketing Information			
Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC Monograph Drug	M012	07/08/2021	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 10/2023 The Procter & Gamble Manufacturing Company