

**VICKS FORMULA 44DM HONEY COUGH AND CHEST CONGESTION-**  
**dextromethorphan hbr, guaifenesin liquid**  
**The Procter & Gamble Manufacturing Company**

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**Vicks ® Formula 44 ™ DM Cough & Congestion Honey Flavored Liquid**  
**Drug Facts**

**Active ingredients (in each 15 mL)**

Dextromethorphan HBr 10 mg  
Guaifenesin 200 mg

**Purpose**

Cough suppressant  
Expectorant

**Uses**

- temporarily relieves common cold symptoms:
- cough due to minor throat & bronchial irritation
- 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**

- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis or emphysema
- a sodium-restricted diet

**Stop use and ask a doctor if**

- 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 6 doses per 24 hours

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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

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**Other information**

- **each 15 mL contains:** sodium 46 mg
- store at no greater than 25°C

**Inactive ingredients**

anhydrous citric acid, D&C Yellow No. 10, FD&C Green No. 3, FD&C Red No. 40, FD&C Yellow No. 6, flavor, glycerin, propylene glycol, saccharin sodium, sodium benzoate, sodium chloride, sorbitol, sucralose, trisodium citrate dihydrate, water, xanthan gum

**Questions?**

**1-800-362-1683**

**TAMPER EVIDENT: DO NOT USE IF PRINTED SHRINKBAND ON BOTTLE IS  
BROKEN OR MISSING.**

**PRINCIPAL DISPLAY PANEL - 354 ml Bottle Label**

**VICKS®**

**FORMULA 44DM**

**HONEY COUGH & CONGESTION**

Dextromethorphan HBr

Guaifenesin

Cough

Chest Congestion, Thins & Loosens Mucus

12 FL OZ (354 mL)



**Drug Facts (continued)**

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children under 4 years	do not use

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← **PEEL BACK FOR DRUG FACTS**

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CINCINNATI, OH 45202  
Patents: [www.pg.com/patents](http://www.pg.com/patents)

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**P&G**  
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## VICKS FORMULA 44DM HONEY COUGH AND CHEST CONGESTION

dextromethorphan hbr, guaifenesin liquid

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84126-355
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 30 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 30 mL

Inactive Ingredients	
Ingredient Name	Strength
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SACCHARIN SODIUM</b> (UNII: SB8ZUX40TY)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>CITRIC ACID</b> (UNII: 2968PHW8QP)	
<b>D&amp;C YELLOW NO. 10</b> (UNII: 35SW5USQ3G)	
<b>FD&amp;C GREEN NO. 3</b> (UNII: 3P3ONR6O1S)	
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>TRISODIUM CITRATE DIHYDRATE</b> (UNII: B22547B95K)	

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

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84126-355-12	354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/01/2025	

**Labeler** - The Procter & Gamble Manufacturing Company (004238200)