

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

VICKS® FORMULA 44™ DM
COUGH & CHEST CONGESTION

Drug Facts

Active ingredient (in each 15 mL)

Dextromethorphan HBr 10 mg

Guaifenesin 200 mg

Purpose

Cough suppressant

Expectorant

Uses

- temporarily relieves cough due to minor throat and bronchial irritation associated with a cold
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus

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

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

Stop use and ask a doctor if

cough persists for more than 1 week, tends to recur, or is accompanied by fever, rash, or persistent headache. A persistent cough may be a sign 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

adults and children 12 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

Other information

- **each 15 mL contains:**sodium 45 mg
- store at no greater than 25°C

Inactive ingredients

citric acid, FD&C Red No. 40, flavor, glycerin, propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium chloride, sodium citrate, sorbitol, sucralose, xanthan gum

Questions?

1-800-362-1683

TAMPER EVIDENT: Do not use if printed shrinkband is broken or missing.

**DIST. BY PROCTER & GAMBLE,
CINCINNATI OH 45202**

PRINCIPAL DISPLAY PANEL - 354 ml Bottle Label

VICKS®

FORMULA 44™ DM

COUGH &

CHEST CONGESTION

Dextromethorphan HBr, Guaifenesin

Cough

Chest Congestion, Thins & Loosens Mucus

Non-Drowsy

12 FL OZ (354 ml)



Drug Facts (continued)

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← PEEL BACK FOR DRUG FACTS

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DIST. BY PROCTER & GAMBLE, CINCINNATI OH 45202
Patents: www.pg.com/patents
www.vicks.com

PARENTS:
Learn and use medicine safely
www.StopMedicineAbuse.org

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VICKS FORMULA 44DM COUGH AND CHEST CONGESTION

dextromethorphan hbr, guaifenesin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69423-810
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 15 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SODIUM CHLORIDE (UNII: 451W471Q8X)	

SORBITOL (UNII: 506T60A25R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C00X)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

Product Characteristics

Color	red	Score	
Shape		Size	
Flavor	BERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-810-12	354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/21/2020	
2	NDC:69423-810-24	2 in 1 CELLO PACK	07/01/2025	
2		354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/21/2020	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 12/2025

The Procter & Gamble Manufacturing Company