

**ADULT COUGH AND CHEST CONGESTION RELIEF DM NON DROWSY-
dextromethorphan hbr, guaifenesin liquid
Safeway, Inc.**

Drug Facts

Active ingredients (in each 10 mL)

Dextromethorphan HBr 20 mg

Guaifenesin 200 mg

Purposes

Cough suppressant

Expectorant

Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes

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)
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema

Stop use and ask a doctor if

cough lasts more than 7 days, comes back, or is accompanied by 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 (1-800-222-

1222) right away.

Directions

- do not take more than 6 doses in any 24-hours period
- measure only with dosing cup provided. Do not use any other dosing device
- keep dosing cup with product
- mL = milliliter
- this adult product is not intended for use in children under 12 years of age
- adults and children 12 years and over: 10 mL every 4 hours
- children under 12 years: do not use

Other information

- store between 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

anhydrous citric acid, FD&C red #40, flavor, glucose, glycerin, high fructose corn syrup, menthol, purified water, saccharin sodium, sodium benzoate

Questions or comments?

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

Principal Display Panel

COMPARE TO Robitussin® Cough + Chest Congestion DM active ingredients**

TUSSIN

COUGH & CHEST CONGESTION DM

FOR AGES 12 AND OVER

DEXTROMETHORPHAN HBr, 20 mg

Cough Suppressant

GUAIFENESIN 200 mg

Expectorant

Alcohol-Free

Relieves

- Cough
- Mucus

**This product is not manufactured or distributed by Haleon group of companies distributors of Robitussin® Cough + Chest Congestion DM.

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF PRINTED SAFETY

SEAL AROUND BOTTLE OR UNDER CAP IS BROKEN OR MISSING.

DISTRIBUTED BY:

BETTER LIVING BRANDS LLC

P.O BOX 99,

PLEASANTON, CA 94566-0009

Package Label



Max ID: FC009078~F CAD: PA220029_61-DI000472	Cust. ID: FC009078~F Customer: PLD	Emboss / Deboss ID N/A	Proof Created: 08/18/2025
Dieline:			
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Customer Approval:	Date:		

Signature Care ADULT COUGH AND CHEST CONGESTION RELIEF DM

ADULT COUGH AND CHEST CONGESTION RELIEF DM NON DROWSY

dextromethorphan hbr, guaifenesin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-385
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 10 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
MENTHOL (UNII: L7T10EIP3A)	
GLUCOSE OXIDASE (UNII: 0T8392U5N1)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-385-04	1 in 1 BOX	06/30/2014	
1		118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:21130-385-08	1 in 1 BOX	06/30/2014	
2		237 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	06/30/2014	

