COUGH CONTROL DM MAX- dextromethorphan hbr, guaifenesin liquid
Select Brand Dist.

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.

Drug Facts

Active ingredients (in each 5 mL, 1 teaspoon)

Dextromethorphan HBr, USP 10 mg

Guaifenesin, USP 200 mg

Purpose

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

When using this product

- **do not use more than directed**

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 this and all drugs out of the reach of children.

In case of accidental overdose, seek professional assistance or contact a Poison Control Center (1-800-222-1222) immediately.

Directions

- **shake well before using**
- do not exceed 6 doses in a 24-hour period
- this adult product is not intended for use in children under 12 years of age
- tsp = teaspoon, mL = milliliter

age	dose
adults and children 12 years and over	2 teaspoonfuls (10 mL) every 4 hours
children under 12 years	do not use

Other information

- each teaspoon contains: **sodium 5 mg**
- store at room temperature
- alcohol-free
- dosage cup provided

Inactive ingredients

anhydrous citric acid, carboxymethylcellulose sodium, flavors, glycerin, high fructose corn syrup, menthol, microcrystalline cellulose, polyethylene glycol, povidone, propylene glycol, purified water, red 33, red 40, saccharin sodium, sodium benzoate, sorbitol, xanthan gum

Principal Display Panel

COUGH CONTROL DM MAX

Cough Suppressant (Dextromethorphan HBr)

Expectorant (Guaifenesin)

Maximum Strength for MUCUS RELIEF

NON-DROWSY

- Controls Coughs
- Relieves Mucus/Chest Congestion

ALCOHOL-FREE

For Adults

*Compare to the active ingredients in Robitussin® DM Max

SEE NEW DOSING INFORMATION

FL OZ (mL)

*This product is not manufactured or distributed by Wyeth Consumer Healthcare, distributors of Robitussin® DM Max.

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL ON THE BOTTLE IS BROKEN OR MISSING.

Distributed by: SELECT BRAND DISTRIBUTORS

Pine Bluff, AR 71603 USA AC (870) 535-3635

MADE IN THE USA

Package Label

Select Brand Cough Control DM Max Liquid

COUGH CONTROL DM MAX

dextromethorphan hbr, guaifenesin liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:15127-864
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 5 mL

GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 5 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
GLYCERIN (UNII: PDC6A3C0OX)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
MENTHOL (UNII: L7T10EIP3A)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL (UNII: 506T60A25R)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:15127-864-04	1 in 1 BOX	09/08/2010	12/31/2020
1		118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:15127-864-08	1 in 1 BOX	09/08/2010	12/31/2020
2		236 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 FINAL	part341	09/08/2010	12/31/2020

Labeler - Select Brand Dist. (012578514)

Revised: 8/2019 Select Brand Dist.