MAXIMUM STRENGTH ADULT COUGH PLUS CHEST CONGESTION DM-
dextromethorphan hbr, guaifenesin solution
FAMILY DOLLAR SERVICES INC

Maximum Strength Adult Cough+Chest Congestion DM Drug Facts

Active ingredients (in each 20 mL)

Dextromethorphan HBr, USP 20 mg

Guaifenesin, USP 400 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 for more than 7 days, comes back, 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.

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Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Directions

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided
- keep dosing cup with product
- mL = milliliter
- this adult product is not intended for use in children under 12 years of age

age	dose
adults and children 12 years and over	20 mL every 4 hours
children under 12 years	do not use

Other information

- **each 20 mL contains:**sodium 7 mg
- store at room temperature. Do not refrigerate.

Inactive ingredients

anhydrous citric acid, carboxymethylcellulose sodium, edetate disodium, FD&C Red No. 40, flavor, menthol, microcrystalline cellulose, povidone, propylene glycol, purified water, potassium citrate, sodium benzoate, sorbitol solution, sucralose, xanthan gum.

Questions or comments?

1-866-467-2748

Package/Label Principal Display Panel

Compare to the active ingredients of Robitussin® Maximum Strength Cough + Chest Congestion DM MAX*

NDC# 55319-740-04

Maximum Strength

Adult

Cough+Chest

Congestion DM

Dextromethorphan HBr (Cough Suppressant)

Guaifenesin (Expectorant)

- Controls cough
- Relieves Chest Congestion
- Thins and Loosens Mucus

Sugar-Free

No Added Alcohol

Use Dosage Cup Included

Natural Raspberry Flavor

For Ages 12 & Over

4 FL OZ (118 mL)

Distributed By:

*This product is not manufactured or distributed by Pfizer, owner of the registered trademark Adult Robitussin ®Maximum Strength Cough + Congestion DM MAX.

MAXIMUM STRENGTH ADULT COUGH PLUS CHEST CONGESTION DM

dextromethorphan hbr, guaifenesin solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55319-740
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 20 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	red	**Score**	
Shape		**Size**	
Flavor	RASPBERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55319-740-04	1 in 1 CARTON	05/08/2023	
1		118 mL in 1 BOTTLE; 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	05/08/2023	

Labeler - FAMILY DOLLAR SERVICES INC (024472631)