MMM - COUGH FORMULA GUAIFENESIN DM- guaifenesin dm liquid
Southern Sales & Services, Inc

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.

MMM Cough Formula
Guaifenesin DM

Active ingredient (in each 5ml tsp.)
Dextromethorphan HBr USP 10 mg.....Cough suppressant

Guaifenesin USP 100mg.....Expectorant

Expectorant, Cough Suppressant

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 you are taking a prescription drug that contains an MAOI drug, ask a doctor or pharmacist before taking this product.

Ask a doctor before use if

you have a cough that lasts or a chronic cough such as occurs with smoking, asthma, chronic bronchitis or emphysema - cough occurs with too much phlegm (mucus)

Stop use and ask a doctor if

cough lasts for more than 7 days, comes back, or occurs with fever, rash, or headache that lasts. These could be signes of a serious condition.

If pregnant or breast-feeding

ask a health professional before use

Keep out of the reach of children

In case of oversdose, get medical help or contact a Poison Control Center right away.

See DOSAGE & ADMINISTRATION

Do not take more than 6 doses in any 24 hour period

adults and children 12 years and over...10ml (2 teaspoonfuls every 4 hours or as directed by doctor)

Children under 12 years...do not use

Store at room temperature - 15°- 30°C (59°- 86°F)

TAMPER-EVIDENT : Do not use if seal under cap is torn, broken or missing.

Propylene glycol, glycerine, citric acid, sucralose, sodium citrate, potassium sorbate, methylparaben, propylparaben, cherry flavor, menthol, FD&C Red#40, purified Water

KEEP OUT OF REACH OF CHILDREN

MMM Cough Formula DM

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MMM - COUGH FORMULA GUAIFENESIN DM

guaifenesin dm liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69822-021
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 5 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 5 mL

Inactive Ingredients	
Ingredient Name	**Strength**
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
CHERRY (UNII: BUC5I9595W)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
MENTHOL (UNII: L7T10EIP3A)	
WATER (UNII: 059QF0KO0R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69822-021-40	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	01/01/2021	

Labeler - Southern Sales & Services, Inc (013114906)

Registrant - Southern Sales & Services, Inc. (013114906)

Establishment

Name	Address	ID/FEI	Business Operations
Southern Sales & Services, Inc.		013114906	label(69822-021)

Revised: 1/2021 Southern Sales & Services, Inc