MUCUS RELIEF DM- dextromethorphan hydrobromide, guaifenesin tablet, extended release Rite Aid Corporation

Rite Aid Corporation Mucus Relief DM Drug Facts

Active ingredients (in each extended-release tablet)

Dextromethorphan HBr 30 mg Guaifenesin 600 mg

Purposes

Cough suppressant

Expectorant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- temporarily relieves:
- cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
- the intensity of coughing
- the impulse to cough to help you get to sleep

Warnings

Do not use

- for children under 12 years of age
- 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 accompanied by too much phlegm (mucus)

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 occurs with fever, rash, or persistent headache. These could be signs of a serious illness.

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. (1-800-222-1222)

Directions

- do not crush, chew, or break tablet
- take with a full glass of water
- this product can be administered without regard for timing of meals
- adults and children 12 years and older:
 - 1 or 2 tablets every 12 hours; not more than 4 tablets in 24 hours
- children under 12 years of age: do not use

Other information

- each tablet contains: magnesium 15 mg
- do not use if printed foil under cap is broken or missing
- store between 20-25°C (68-77°F)

Inactive ingredients

carbomer homopolymer type B, copovidone, D&C yellow #10 aluminum lake, hypromellose, magnesium hydroxide, magnesium stearate, microcrystalline cellulose, silicon dioxide

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

Compare to the active ingredients of Mucinex® DM

FREE FROM

GLUTEN FREE

MUCUS RELIEF DM

GUAIFENESIN and DEXTROMETHORPHAN HYDROBROMIDE EXTENDED-RELEASE TABLETS, 600 mg / 30 mg

EXPECTORANT & COUGH SUPPRESSANT

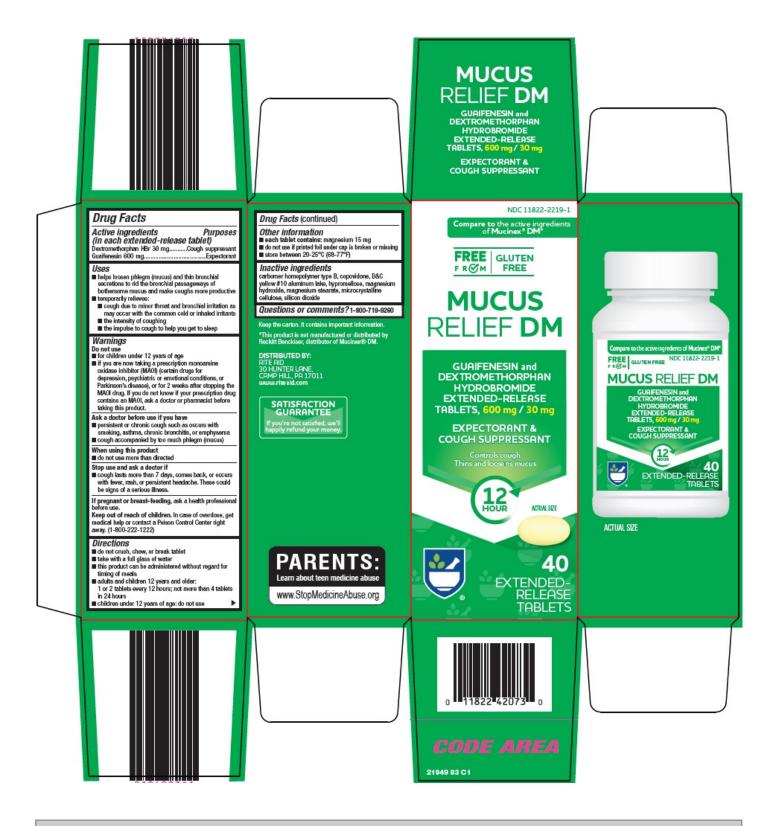
Controls cough

Thins and loosens mucus

12 HOUR

ACTUAL SIZE

40 EXTENDED-RELEASE TABLETS



MUCUS RELIEF DM

dextromethorphan hydrobromide, guaifenesin tablet, extended release

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-2219
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg	
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	600 mg	

Inactive Ingredients		
Ingredient Name	Strength	
CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL OR ALLYL SUCROSE CROSSLINKED) (UNII: K6MOM3T5YL)		
COPOVIDONE K25-31 (UNII: D9C330MD8B)		
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)		
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)		
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		

Product Characteristics				
Color	YELLOW	Score	no score	
Shape	OVAL	Size	16mm	
Flavor		Imprint Code	L219;600	
Contains				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:11822- 2219-1	1 in 1 CARTON	03/16/2023		
1		40 in 1 BOTTLE; Type 0: Not a Combination Product			
2	NDC:11822- 2219-0	1 in 1 CARTON	03/15/2023		
2		20 in 1 BOTTLE; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA207602	03/15/2023	

Labeler - Rite Aid Corporation (014578892)

Revised: 3/2023 Rite Aid Corporation