MUCUS RELIEF DM MAXIMUM STRENGTH- guaifenesin, dextromethorphan hbr tablet Rite Aid Corporation

Drug Facts

Active ingredients (in each extended-release tablet)

Dextromethorphan HBr 60 mg

Guaifenesin 1200 mg

Purpose

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 under12 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 regards for timing of meals
- adults and children 12 years of age and older: 1 tablet every 12 hours; not more than 2 tablets in 24 hours
- children under 12 years of age: do not use

Other information

store between 20^o to 25^oC (68^o to 77^oF)

Inactive ingredients

carbomer, colloidal silicon dioxide, D&C yellow #10 aluminum lake, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, povidone, talc

Questions or comments?

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

Principal Display Panel

Compare to the active ingredients in Maximum Strength Mucinex ® DM* MAXIMUM STRENGTH MUCUS RELIEF DM GUAIFENESIN 1200 mg DEXTROMETHORPHAN HBr 60 mg EXPECTORANT & COUGH SUPPRESSANT Temporarily controls cough Thins & loosens mucus 12 HOUR EXTENDED-RELEASE TABLETS

*This product is not manufactured or distributed Reckitt Benckiser LLC, distributor of Maximum Strength Mucinex ® DM.

TAMPER EVIDENT: DO NOT USE IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

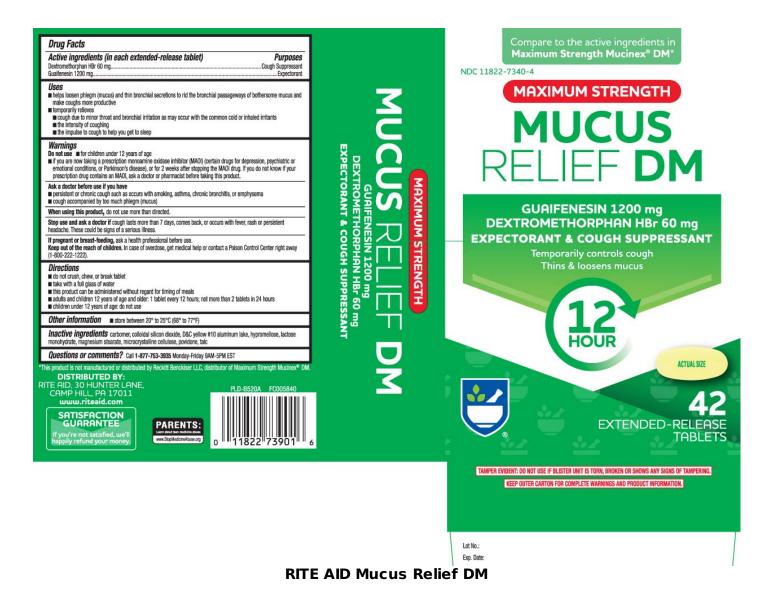
DISTRIBUTED BY:

RITE AID, 30 HUNTER LANE,

CAMP HILL, PA 17011

www.riteaid.com

Package Label



MUCUS REI	IEF DM	ΜΑΧΙΜυ	M STR	ENGT	н			
guaifenesin, dex	tromethorph	an hbr tablet						
Product Infor	mation							
Product Type	HUMAN OTC DRUG Item C				Code (Source) NDC:			2:11822-7340
Route of Admin	istration ORAL							
Active Ingred	ient/Active	Moiety						
	Ingredient Name					Basis of S	gth Strength	
DEXTROMETHORPHAN HYDROBROMIDE (UNII: (DEXTROMETHORPHAN - UNII:7355X3ROTS)						DEXTROMETHORPHAN HYDROBROMIDE		60 mg
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN			- UNII:495W7451VQ) GUAIFENESIN				1200 mg	
Inactive Ingre	edients							
Ingredient Name								Strength
CARBOMER 934 (UNII: Z135WT9208)								
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)								
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)								
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)								
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)								
MAGNESIUM STEARATE (UNII: 70097M6I30) CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)								
POVIDONE (UNII: FZ989GH94E)								
TALC (UNII: 7SEV7J4R1U)								
	11(20)							
Product Chara	acteristics							
Color	low	Score				no score		
Shape	AL	Size				22mm		
Flavor		Imprint Code				AN03	AN039	
Contains								
Packaging								
# Item Code	Pa	ckage Desc	ription		Mar	keting Start	M	larketing End
1 NDC:11822-	42 in 1 CARTO				05/31/2	Date 31/2019		Date
1 7340-4	1 in 1 BLISTER PACK; Type 0: Not a Combination			oination				
	Product							
Marketing	Informat	ion						
Marketing Category		tion Number Citatio		graph	Ma	rketing Start Date	ſ	Marketing End Date
ANDA	ANDA20969		-		05/31	/2019		

Labeler - Rite Aid Corporation (014578892)

Revised: 1/2022

Rite Aid Corporation