# MUCUS RELIEF DM- dextromethorphan hydrobromide, guaifenes in tablet, extended release Meijer Distribution Inc

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## Meijer Distribution, Inc. 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

VALUE SIZE 40 EXTENDED-RELEASE TABLETS

Compare to Mucinex® DM active ingredients

mucus relief DM

Guaifenesin 600 mg & Dextromethorphan Hydrobromide 30 mg Extended-Release Tablets

Expectorant | Cough Suppressant

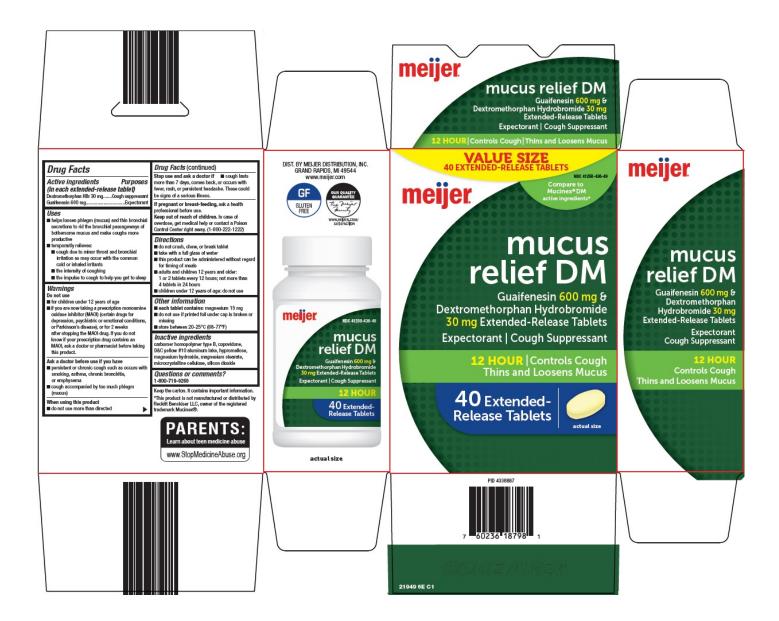
12 HOUR

Controls Cough

Thins and Loosens Mucus

40 Extended-Release Tablets

actual size



#### **MUCUS RELIEF DM**

dextromethorphan hydrobromide, guaifenesin tablet, extended release

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41250-436
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9 D2RTI9 KYH) (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: D9 C330 MD8 B)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM HYDRO XIDE (UNII: NBZ3Q Y004S)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U)	
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)	

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

I	Packaging				
#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>	
1	NDC:41250-436-01	1 in 1 CARTON	06/02/2020		
1		20 in 1 BOTTLE; Type 0: Not a Combination Product			
2	NDC:41250-436-49	1 in 1 CARTON	06/02/2020		
2		40 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	06/02/2020	

# Labeler - Meijer Distribution Inc (006959555)

Revised: 6/2020 Meijer Distribution Inc