MUCUS RELIEF DM - guaifenesin and dextromethorphan hbr tablet, extended release

Wal-Mart Stores, Inc.

Drug Facts

Active Ingredients

(in each extended-release tablet)

Dextromethorphan Hydrobromide USP 30 mg Guaifenesin USP 600 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 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 (1-800-222-1222) right away.

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

- tamper evident: do not use if seal over bottle opening is broken or missing
- store at 20° to 25°C (68° to 77°F)

Inactive ingredients

colloidal silicon dioxide, hypromellose, magnesium stearate, microcrystalline cellulose, povidone, pregelatinized starch (maize)

Questions?

call **1-888-287-1915** You may also report side effects to this phone number.

DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716

PRODUCT OF INDIA

*This product is not manufactured or distributed by Reckitt Benckiser, owner of the registered trademark Mucinex®DM.

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 600 mg/30 mg (20 Tablet Carton Label)

NDC 49035-960-73 equateTM
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

20 EXTENDED-RELEASE TABLETS

62



PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 600 mg/30 mg (20 Tablet Carton Label)

NDC 49035-960-73

equateTM

Mucus Relief DM

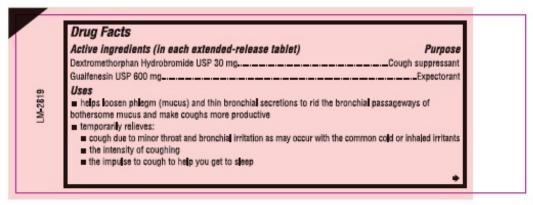
Guaifenesin 600 mg / Dextromethorphan Hydrobromide 30 mg Extended-Release Tablets Expectorant/ Cough Suppressant

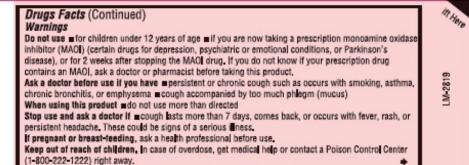
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MUCUS RELIEF DM

guaifenesin and dextromethorphan hbr tablet, extended release

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-960	
Route of Administration	ORAL			

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

Inactive Ingredients				
Ingredient Name	Strength			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
POVIDONE K90 (UNII: RDH86HJV5Z)				
POVIDONE K25 (UNII: K0KQV10C35)				
STARCH, CORN (UNII: O8232NY3SJ)				

Product Characteristics					
Color	WHITE (white to off-white)	Score	no score		

Shape	OVAL	Size	16mm
Flavor		Imprint Code	X;62
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:49035-960- 73	1 in 1 CARTON	03/17/2017		
1		20 in 1 BOTTLE; Type 0: Not a Combination Product			
2	NDC:49035-960- 12	1 in 1 CARTON	03/17/2017		
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	ANDA206941	03/17/2017		

Labeler - Wal-Mart Stores, Inc. (051957769)

Registrant - Aurohealth LLC (078728447)

Establishment			
Name	Address	ID/FEI	Business Operations
Aurobindo Pharma Limited		650381903	ANALYSIS(49035-960), MANUFACTURE(49035-960)

Revised: 2/2024 Wal-Mart Stores, Inc.