MUCUS RELIEF DM EXTENDED RELEASE CAPLETS- guaifenesin, dextromethorphan hbr tablet EQUATE (Wal-Mart Stores, Inc.) (see also WAL-MART INC)

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° to 25°C (68° to 77°F)

Inactive ingredients

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

Questions or comments?

1-888-287-1915

Principal Display Panel

Compare to Maximum Strength Mucinex® DM active ingredients*

Mucus Relief DM

Guaifenesin 1200 mg/

Dextromethorphan HBr 60 mg

Extended-Release Tablets

MAXIMUM STRENGTH

Expectorant/Cough Suppressant

- Controls Cough
- Thins & Loosens Mucus

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 PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

DISTRIBUTED BY: Walmart Inc.,

Bentonville, AR 72716

Package Label



EQUATE Mucus Relief DM

Lot No.: Exp. Date:

MUCUS RELIEF DM EXTENDED RELEASE CAPLETS

quaifenesin, dextromethorphan hbr tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-824
Poute of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	60 mg	
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	1200 mg	

Inactive Ingredients				
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: FZ 989GH94E)				
TALC (UNII: 7SEV7J4R1U)				

Product Characteristics				
Color	yellow	Score	no score	
Shape	OVAL	Size	22mm	
Flavor		Imprint Code	AN039	
Contains				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:49035- 824-28	1 in 1 BOX	11/29/2019		
1		28 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			

Marketing I	Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA209692	11/29/2019		

Revised: 12/2022