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

Strategic Sourcing Services LLC

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?

Call 833-358-6431 Monday to Friday 9:00am to 7:00pm EST

Principal Display Panel

COMPARE TO MAXIMUM STRENGTH MUCINEX® DM ACTIVE INGREDIENTS*

Mucus Relief DM

Guaifenesin 1200 mg

Dextromethorphan HBr 60 mg

EXPECTORANT

COUGH SUPPRESSANT

12 hour relief of:

- Controls Cough
- Thins and 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 BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGN OF TAMPERING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Distributed by: McKesson Corp.,

via Strategic Sourcing Services LLC.

Memphis, TN 38141

Package Label



FOSTER & THRIVE Maximum Strength Mucus Relief DM

MUCUS RELIEF DM

guaifenesin, dextromethorphan hbr tablet, extended release

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70677-1049(NDC:65162-039)
Route 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 (light 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:70677- 1049-1	14 in 1 CARTON	03/01/2023	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA209692	03/01/2023		

Revised: 4/2023