MUCUS RELIEF DM MAX MAXIMUM STRENGTH- dextromethorphan hbr, guaifenesin tablet
TARGET Corporation

Drug Facts

Active ingredients (in each extended-release tablets)

Dextromethorphan HBr 60 mg

Guaifenesin 1200 mg

Purpose

Cough suppressant

Expectorant

Uses

- help 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
- adult 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 20º to 25º C (68º to 77ºF)

Inactive ingredients

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

Questions?

Call 1-800-910-6874

Principal Display Panel

Compare to active ingredients in Maximum Strength Mucinex® DM*

Maximum Strength

mucus relief DM

guaifenesin 1,200 mg

expectorant

dextromethorphan HBr 60 mg

cough suppressant

controls cough

thins and loosens mucus

EXTENDED-RELEASE TABLETS

*This product is not manufactured or distributed by Reckitt Benckiser, 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 Target Corporation

Minneapolis, MN 55403

Package Label

TARGET Maximum Strength Mucus Relief DM

MUCUS RELIEF DM MAX MAXIMUM STRENGTH

dextromethorphan hbr, guaifenesin tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-834
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)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POVIDONE (UNII: FZ989GH94E)	
TALC (UNII: 7SEV7J4R1U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11673-834-28	28 in 1 CARTON	01/01/2019	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11673-834-14	14 in 1 CARTON	01/01/2019	
2		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	01/01/2019	

Labeler - TARGET Corporation (006961700)

Revised: 10/2019 TARGET Corporation