MUCUS RELIEF ER DM EXTENDED RELEASE- guaifenesin and dextromethorphan hydrobromide tablet, extended release
KROGER COMPANY

Mucus Relief ER DM
Extended Release

Drug Facts

Active ingredients (in each extended-release tablet)	*Purposes*
Dextromethorphan HBr 30 mg	Cough suppressant
Guaifenesin 600 mg	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

- Tamper evident: Do not use if carton is open or if printed seal on blister is broken or missing.
- store between 20-25°C (68-77°F)

Inactive ingredients

carbomer homopolymer type B; hypromellose, USP; magnesium stearate, NF; microcrystalline cellulose, NF; sodium starch glycolate, NF

Questions?

1-800-632-6900

You may also report side effects to this phone number.

DISTRIBUTED BY THE KROGER CO.
CINCINNATI, OHIO 45202

PRINCIPAL DISPLAY PANEL - 20 Tablet Blister Pack Carton

COMPARE TO the active ingredients of
MUCINEX® DM*
See back panel

NDC 30142-991-20

Kroger®

OUR PHARMACIST RECOMMENDED

Mucus Relief
ER DM

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

EXPECTORANT &
COUGH SUPPRESSANT

12
HOUR

- Controls Cough
- Thins and Loosens Mucus
- Immediate & Extended Release

actual
size

20 EXTENDED-RELEASE
TABLETS

MUCUS RELIEF ER DM EXTENDED RELEASE

guaifenesin and dextromethorphan hydrobromide tablet, extended release

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:30142-991
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 (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	Dextromethorphan Hydrobromide	30 mg

Inactive Ingredients

Ingredient Name	Strength
CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: HHT01ZNK31)	

hypromellose, unspecified (UNII: 3NXW29V3WO)	
magnesium stearate (UNII: 70097M6I30)	
microcrystalline cellulose (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	WHITE	**Score**	no score
Shape	OVAL	**Size**	16mm
Flavor		**Imprint Code**	xeunciM;600
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30142-991-20	1 in 1 CARTON	05/30/2019	
1		20 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:30142-991-40	2 in 1 CARTON	05/30/2019	
2		40 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
NDA	NDA021620	05/30/2019	

Labeler - KROGER COMPANY (006999528)

Establishment

Name	Address	ID/FEI	Business Operations
RECKITT BENCKISER HEALTHCARE INTERNATIONAL LTD		230780363	MANUFACTURE(30142-991) , LABEL(30142-991)

Revised: 2/2020 KROGER COMPANY