MUCINEX DM - guaifenes in and dextromethorphan hydrobromide tablet, extended release Physicians Total Care, Inc.

Mucinex®DM

600 mg guaifenesin & 30 mg dextromethorphan HBr extended-release bi-layer tablets EXPECTORANT & COUGH SUPPRESSANT

Drug Facts

Active ingredients (in each extended-release bi-layer tablet)	Purpose
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.

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 at 20-25°C (68-77°F)

Inactive ingredients

carbomer 934P, NF; D&C yellow #10 aluminum lake; hypromellose, USP; magnesium stearate, NF; microcrystalline cellulose, NF; sodium starch glycolate, NF

Distributed by: Reckitt Benckiser Inc. Parsippany, NJ 07054-0224 ©RBI 2009

Relabeling of "Additional Barcode" by: Physicians Total Care, Inc. Tulsa, OK 74146

PRINCIPAL DISPLAY PANEL - 600 mg Carton

NDC 54868-5853-0

Mucinex®DM

600 mg guaifenes in & 30 mg dextromethorphan HBr extended-releas e bi-layer tablets

EXPECTORANT & COUGH SUPPRESSANT

12 HOUR

NEW LOOK-SAME RELIEF

Thins And Loosens Mucus

Controls Cough

20 EXTENDED-RELEASE BI-LAYER TABLETS



MUCINEX DM

guaifenesin and dextromethorphan hydrobromide tablet, extended release

Produc	t Infor	mation
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:54868-5853(NDC:63824-056)

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		
CARBOMER 934 (UNII: Z135WT9208)			
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)			
HYPROMELLOSE (UNII: 3NXW29 V3WO)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
CELLULOSE, MICRO CRYSTALLINE (UNII: OP1R32D61U)			
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)			

Product Characteristics			
Color	white (yellow and white)	Score	no score
Shape	OVAL	Size	16 mm
Flavor		Imprint Code	Mucinex;600
Contains			

racka	ging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:	54868-5853-0	2 in 1 CARTON		
1		10 in 1 BLISTER PACK		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA021620	01/23/2008	

Labeler - Physicians Total Care, Inc. (194123980)

Establishment			
Name	Address	ID/FEI	Business Operations
Physicians Total Care, Inc.		194123980	relabel

Revised: 12/2009 Physicians Total Care, Inc.