HEALTHMART MUCUS RELIEF FM- acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl tablet Mckesson

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Mucus Relief FM

Active ingredients (in each caplet)

Acetaminophen 325 mg Dextromethorphan HBr 10 mg Guaifenesin 200 mg Phenylephrine HCl 5 mg

Purpose

Pain reliever/Fever reducer

Cough Suppressant

Expectorant

Nasal decongestant

Uses

- temporarily relieves these common cold and flu symptoms:
- nasal congestion
- cough
- headache
- fever
- minor aches and pains
- sinus congestion and pressure
- temporarily promotes nasal and/or sinus drainage
- helps loosen (mucus) and thin bronchial secretions to drain bronchial tubes and make coughs more productive.

Warnings

Liver warning: This product contains acetaminophen. severe liver damage may occur if you take:

- more than 12 caplets in 24 hours, which is maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks daily while using this product

Do not use

- with any other drug containing acetaminophen (prescription or non prescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- 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

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis or emphysema
- cough that occurs with too much phlegm (mucus)

Ask a doctor or pharmacist before use if

you are taking the blood thinning drug warfarin

When using this product

do not use more than directed

Stop use and ask a doctor if

- nervousness, dizziness or sleeplessness occur
- pain, nasal congestion or cough gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- cough comes back, or occurs with rash or persistent headache. These could be signs of a serious condition.

If pregnant or breast-feeding,

ask a health professional before use. Keep out of reach of children.

Overdose warning

Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a poison control center right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

- do not take more than directed (see overdose warning)
- do not take more than 12 caplets in any 24 hour period
- Adults and children 12 years and older: take 2 caplets every 4 hours.
- Children under 12 years of age: Do not use

Other information

- tamper evident: do not use if individual blister units are borken or open
- store between 15-30°C(59-86°F)

Inactive ingredients

Colloidal silicon dioxide, croscaramellose sodium, crospovidone, FD&C red 40 aluminum lake, FD&C

yellow 6 aluminum lake, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, stearic acid, talc, titanium dioxide. May contain methacrylic acid copolymer, sodium bicarbonate

PDP

Health Mart

compare to Mucinex Fast-MaxTM severe congestion & cold caplets active ingredients

for ages 12+

mucus relief FM

adult severe congestion & cold

acetaminophen.....pain reliever/Fever reducer

dextromethorphan HBr.....cough Suppressant

guaifenesin....expectorant

phenylephrine HCl.....nasal congestion

soothing relief for headache, fever, cough, nasal and chest congestion

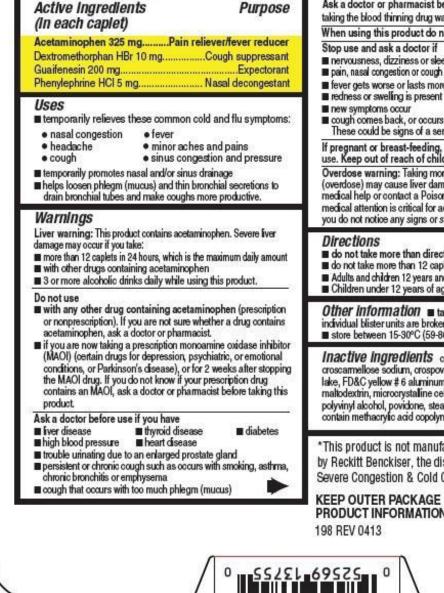
thins and loosens mucus

20 Caplets



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Drug Facts



HEALTHMART MUCUS RELIEF FM

acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl tablet

MFR# 53041 REV 0513

Drug Facts (continued)

Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin.

When using this product do not use more than directed.

Stop use and ask a doctor if

- nervousness, dizziness or sleeplessness occur
- pain, nasal congestion or cough gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days

- cough comes back, or occurs with rash or persistent headache. These could be signs of a serious condition.

If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children.

Overdose warning: Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

- do not take more than directed (see Overdose warning)
- do not take more than 12 caplets in any 24-hour period Adults and children 12 years and older: take 2 caplets every 4 hours.
- Children under 12 years of age: do not use

Other information a tamper evident: do not use if individual blister units are broken or open store between 15-30°C (59-86°F)

Inactive ingredients colloidal silicon dioxide, croscarmellose sodium, crospovidone, FD&C red # 40 aluminum lake, FD&C yellow # 6 aluminum lake, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, stearic acid, talc, titanium dioxide. May contain methacrylic acid copolymer, sodium bicarbonate.

*This product is not manufactured or distributed by Reckitt Benckiser, the distributor of Mucinex® Fast-Max™ Severe Congestion & Cold Caplets.

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62011-0218
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9 D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg
PHENYLEPHRINE HYDRO CHLO RIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients	
Ingredient Name	Strength
COLLOIDAL SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
CROSPOVIDONE (UNII: 68401960MK)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO.6 (UNII: H77VEI93A8)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTO DEXTRIN (UNII: 7CVR7L4A2D)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
PO VIDO NE (UNII: FZ989GH94E)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)	

Other ingreatents					
Ingredient Kind	Ingredient Name Quan			Quantity	
May contain	METHACRYLIC ACID (UNII: 1CS02G8656)				
May contain	SODIUM BICARBONA	TE (UNII: 8	MDF5V39QO)		
Product Characteristics					
Color	red	Score		no score	
Shape	capsule	Size		19 mm	
Flavor		Imp rint O	Code	G198	
Contains					
Packaging					
# Item Code	Package Descriptio	n	Marketing Start Date	Marketi	ng End Date

1 NDC:62011-0218-1	2 in 1 CARTON		
1	10 in 1 BLISTER PACK		
Marketing Info	rmation		
Marketing Info Marketing Category	r mation Application Number or Monograp	h Citation Marketing Start Dat	e Marketing End Date
Marketing Info Marketing Category OTC monograph final		h Citation Marketing Start Dat 04/04/2013	e Marketing End Date

Labeler - Mckesson (177667227)

Revised: 4/2014

Mckesson