MAXIMUM STRENGTH MUCINEX SINUS-MAX PRESSURE, PAIN AND COUGH-
acetaminophen, guaifenesin tablet, film coated
RB Health (US) LLC

Active ingredients
(in each caplet)

Acetaminophen 325 mg
Guaifenesin 200 mg

Uses
■ temporarily relieves these common cold and flu symptoms:
■ minor aches and pains ■ headache
■ sore throat
■ helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
■ temporarily reduces fever

Warnings
Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take:
■ more than 12 caplets in 24 hours, which is the maximum daily amount for this product
■ with other drugs containing acetaminophen
■ 3 or more alcoholic drinks daily while using this product

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:
■ skin reddening ■ blisters ■ rash
If a skin reaction occurs, stop use and seek medical help right away.

Sore throat warning: If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

Do not use
■ with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.

Ask a doctor before use if you have
■ liver disease
■ 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
■ pain 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 headache that lasts. These could be signs of a serious condition.

If pregnant or breast-feeding, ask a health professional before use.

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 of age and over: take 2 caplets every 4 hours
■ children under 12 years of age: do not use

Other information
■ store at 20-25°C (68-77°F)

Inactive ingredients
croscarmellose sodium, FD&C red no. 40 aluminum lake, FD&C yellow no. 6 aluminum lake, magnesium stearate, microcrystalline

cellulose, polyethylene glycol, polyvinyl alcohol, povidone, talc, titanium dioxide

Questions? 1-866-MUCINEX
(1-866-682-4639)

Acetaminophen 325 mg ..... Pain reliever/
fever reducer
Guaifenesin 200 mg..... Expectorant

Keep out of reach of children.

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acetaminophen, guaifenesin tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72854-208
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg

Inactive Ingredients

Ingredient Name	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
TALC (UNII: 7SEV7J4R1U)	
FD&C RED NO. 40 ALUMINUM LAKE (UNII: 6T47AS764T)	
FD&C YELLOW NO. 6 ALUMINUM LAKE (UNII: GYP6Z2JR6Q)	

Product Characteristics

Color	red	**Score**	no score
Shape	OVAL	**Size**	20mm
Flavor		**Imprint Code**	VVV;MSC
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72854-208-10	1 in 1 CARTON	05/01/2025	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:72854-208-20	2 in 1 CARTON	05/01/2025	
2		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:72854-208-02	2 in 1 POUCH; Type 0: Not a Combination Product	05/01/2025	
4	NDC:72854-208-08	4 in 1 CARTON	05/01/2025	
4		2 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	05/01/2025	

Labeler - RB Health (US) LLC (081049410)

Revised: 5/2025 RB Health (US) LLC