MUCINEX RAPID CLEAR PAIN, HEADACHE, AND MUCUS CONGESTION AND SINUS MAX CLEAR AND COOL- oxymetazoline hydrochloride, acetaminophen and guaifenesin RB Health (US) LLC

MUCINEX® RAPID+CLEAR Pain Headache & Mucus Congestion + MUCINEX® SINUS-MAX® Severe Nasal Congestion Relief CLEAR & COOL Nasal Spray

Active ingredient Purpose

Oxymetazoline hydrochloride 0.05%......Nasal decongestant

Active ingredients (in each capsule) Purposes

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

Uses

- temporarily relieves nasal congestion due to:
- a cold hay fever or other upper respiratory allergies
- promotes nasal and/or sinus drainage; temporarily relieves sinus congestion and pressure
- helps clear nasal passages; shrinks swollen membranes

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

Ask a doctor before use if you have

- heart disease
 high blood pressure
- thyroid disease
 diabetes
- difficulty in urination due to enlargement of the prostate gland

When using this product

- do not exceed recommended dosage
- do not use this product for more than 3 days. Use only as directed. Frequent or prolonged use

may cause nasal congestion to recur or worsen.

■ this product may cause temporary discomfort such as burning, stinging, sneezing, or an

increase in nasal discharge

■ the use of this container by more than one person may spread infection

Warnings

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

- more than 12 capsules 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.

Directions

■ adults and children 6 to under 12 years of age (with adult supervision): 2 or 3 sprays in each

nostril, not more often than every 10 to 12 hours. Do not exceed 2 doses in any 24-hour period.

■ children under 6 years of age: consult a doctor.

Shake well before use.

To open, hold by the white grips then squeeze, push down firmly and turn cap counterclockwise.

Before using for the first time, remove the protective cap from the tip and prime metered pump

by depressing firmly several times. To spray, hold bottle with thumb at the base and nozzle

between first and second fingers. Without tilting head, insert nozzle into nostril. Fully depress

pump all the way down with a firm even stroke and sniff deeply. Wipe nozzle clean after use.

To close, turn cap clockwise.

DO NOT DISCARD CAP.

Directions

- do not take more than directed (see Overdose warning)
- do not take more than 12 capsules in any 24-hour period
- adults and children 12 years of age and over: take 2 capsules every 4 hours
- children under 12 years of age: do not use

Other information

- store at 20-25°C (68-77°F)
- avoid excessive heat

Inactive ingredients

benzalkonium chloride, benzododecinium chloride, camphor, cetalkonium chloride, colloidal

silicon dioxide, edetate disodium, eucalyptol, glycine, linoleic acid, linolenic acid, menthol, myristalkonium chloride, myristic acid, oleic acid, palmitic acid, palmitoleic acid, polyethylene

glycol, polysorbate 80, propylene glycol, purified water, sodium carbonate, sodium chloride,

sodium hydroxide, stearic acid

Inactive ingredients

FD&C blue no.1, gelatin, glycerin, hypromellose, lecithin, light mineral oil, polyethylene glycol,

povidone, propylene glycol, purified water, sorbitol sorbitan solution, titanium dioxide Keep out of reach of Children

Purpose

Oxymetazoline hydrochloride 0.05%......Nasal decongestant

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



Severe Nasal Congestion Relief Clear & Cool Nasal Spray Pain, Headache & Mucus Congestion Drug Facts **Drug Facts** Active ingredient Purpose Active ingredients (in each capsule) Purposes Oxymetazoline hydrochloride 0.05%.Nasal decongestant Guaifenesin 200 mg. temporarily relieves nasal congestion due to: ■ temporarily relieves these common cold and flu symptoms: a cold hay fever or other upper respiratory allergies promotes nasal and/or sinus drainage; temporarily relieves sinus congestion and pressure ■ minor aches and pains ■ headache ■ sore throat helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways helps clear nasal passages; shrinks swollen membranes of bothersome mucus and make coughs more productive temporarily reduces fever Warnings Warnings Ask a doctor before use if you have ■ heart disease ■ high blood pressure ■ thyroid disease ■ diabetes Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take: more than 12 capsules 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 difficulty in urination due to enlargement of the prostate gland When using this product Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include: do not exceed recommended dosage ■ skin reddening ■ blisters ■ rash ■ do not use this product for more than 3 days. Use only as directed. Frequent or prolonged use If a skin reaction occurs, stop use and seek medical help right away. may cause nasal congestion to recur or worsen. Sore throat warning: If sore throat is severe, persists for more than 2 days, is accompanied or this product may cause temporary discomfort such as burning, stinging, sneezing, or an followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly. increase in nasal discharge Do not use the use of this container by more than one person may spread infection with any other drug containing acetaminophen (prescription or nonprescription). If you are not Stop use and ask a doctor if symptoms persist sure whether a drug contains acetaminophen, ask a doctor or pharmacist. If pregnant or breast-feeding, ask a health professional before use. Ask a doctor before use if you have liver disease Keep out of reach of children. If swallowed, get medical help or contact a Poison Control persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema cough that occurs with too much phlegm (mucus) Directions Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin adults and children 6 to under 12 years of age (with adult supervision): 2 or 3 sprays in each nostril, not more often than every 10 to 12 hours. Do not exceed 2 doses in any 24-hour period. When using this product do not use more than directed children under 6 years of age: consult a doctor. 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 Shake well before use. cough comes back or occurs with rash or headache that lasts. These could be signs of a To open, hold by the white grips then squeeze, push down firmly and turn cap counterclockwise. Before using for the first time, remove the protective cap from the tip and prime metered pump serious condition. new symptoms occur by depressing firmly several times. To spray, hold bottle with thumb at the base and nozzle If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of between first and second fingers. Without tilting head, insert nozzle into nostril. Fully depress children. Overdose warning: Taking more than the recommended dose (overdose) may cause pump all the way down with a firm even stroke and sniff deeply. Wipe nozzle clean after use. liver damage. In case of overdose, get medical help or contact a Poison Control Center right To close, turn cap clockwise. away. Quick medical attention is critical for adults as well as for children even if you do not DO NOT DISCARD CAP. notice any signs or symptoms. Other information Directions store at 20-25°C (68-77°F) do not take more than directed (see Overdose warning) do not take more than 12 capsules in any 24-hour period Inactive ingredients adults and children 12 years of age and over: take 2 capsules every 4 hours children under 12 years of age: do not use benzalkonium chloride, benzododecinium chloride, camphor, cetalkonium chloride, colloidal silicon dioxide, edetate disodium, eucalyptoi, glyiche, linoleic acid, linolenic acid, menthol, myristalkonium chloride, myristic acid, oleic acid, palmitic acid, palmitoleic acid, polyethylene Other information ■ store at 20-25°C (68-77°F) avoid excessive heat glýcol, polysorbate 80, propylene glycol, purified water, sodium carbonate, sodium chloride. sodium hydroxide, stearic acid Inactive ingredients FD&C blue no.1, gelatin, glycerin, hypromellose, lecithin, light mineral oil, polyethylene glycol, Questions? 1-866-MUCINEX (1-866-682-4639) povidone, propylene glycol, purified water, sorbitol sorbitan solution, titanium dioxide Questions? 1-866-MUCINEX (1-866-682-4639)

MUCINEX RAPID CLEAR PAIN, HEADACHE, AND MUCUS CONGESTION AND SINUS MAX CLEAR AND COOL

oxymetazoline hydrochloride, acetaminophen and guaifenesin kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:72854-212

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:72854-212- 02	1 in 1 CARTON	05/01/2025			
1		1 in 1 CARTON; Type 0: Not a Combination Product				

Quantity of Parts				
Part #	Package Quantity	Total Product Quantity		
Part 1	0 BLISTER PACK	1		
Part 2	0 BOTTLE, PUMP	1 mL		

Part 1 of 2

MUCINEX SINUS MAX CLEAR AND COOL

acetaminophen and guaifenesin capsule, liquid filled

Product Information

Item Code (Source) NDC:72854-211

Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN 325 mg

Inactive Ingredients			
Ingredient Name	Strength		
GELATIN (UNII: 2G86QN327L)			
GLYCERIN (UNII: PDC6A3C0OX)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
LECITHIN, SUNFLOWER (UNII: 834K0WOS5G)			
SORBITOL SOLUTION (UNII: 8KW3E207O2)			
LIGHT MINERAL OIL (UNII: N6K5787QVP)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
POVIDONE (UNII: FZ 989GH94E)			
WATER (UNII: 059QF0KO0R)			

Product Characteristics				
Color	blue	Score	no score	
Shape	OVAL	Size	23mm	
Flavor		Imprint Code		
Contains				

ı	Packaging				
-	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
:	NDC:72854- 211-16	1 in 1 CARTON			
	L	16 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	
	OTC Monograph Drug	M012	05/01/2025		

Part 2 of 2

MUCINEX SINUS-MAX SEVERE NASAL CONGESTION RELIEF CLEAR AND COOL NASAL

oxymetazoline hydrochloride solution

Product Information

Item Code (Source) NDC:63824-129

Route of Administration NASAL

l	Active Ingredient/Active Moiety				
ı	Ingredient Name	Basis of Strength	Strength		
	OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZ OLINE HYDROCHLORIDE	0.05 g in 100 mL		

Inactive Ingredients			
Ingredient Name	Strength		
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)			
BENZODODECINIUM CHLORIDE (UNII: Y5A751G47H)			
CAMPHOR (NATURAL) (UNII: N20HL7Q941)			

Product Characteristics				
Color		Score		
Shape		Size		
Flavor	MENTHOL	Imprint Code		
Contains				

P	Packaging				
#	# Item Code Package Description		Marketing Start Date	Marketing End Date	
1	NDC:63824- 129-17	1 in 1 CARTON			
1		22 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product			

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

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: 6/2025 RB Health (US) LLC