MUCINEX SINUS MAX CLEAR AND COOL SEVERE CONGESTION- oxymetazoline hci spray RB Health (US) LLC

Mucinex Sinus Max Clear & Cool Severe Congestion

Drug Facts

Active ingredient Oxymetazoline hydrochloride 0.05%

Purpose:

Nasal Decongestant

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

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

Stop use and ask a doctor if symptoms persist

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.

Other information

■ store at 20-25°C (68-77°F)

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

Questions?

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

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

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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Drug Facts (continued)

Wipe nozzle clean after use. To close, turn cap clockwise. DO NOT DISCARD CAP.

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benzalkonium chloride,
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linoleic acid, linolenic acid,
menthol, myristic acid, oleic
acid, palmitic acid, palmitoleic
acid, polyethylene glycol,
polysorbate 80, propylene glycol,
purified water, sodium
carbonate, sodium chloride,
sodium hydroxide, stearic acid

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KEEP CARTON FOR FULL INFORMATION.

Please visit our website

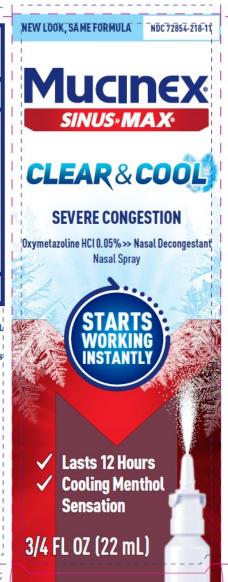
www.mucinex.com

Patents: www.reckitt.com/patents





Dist. by: RB Health (US)
Parsippany NJ 07054-0224
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Made in India



Tamper evident: Do not use if printed seal over cap is torn or missing.

How to Use



1 Open: Hold white grips, squeeze and push down. Turn cap counterclockwise.

Remove cap from tip. Depress several times until mist





3 Apply:
Head upright, insert!
and press down all
the way 2-3 times.
Sniff deeply. Repeat
in other nostril.

After Use: Wipe nozzle clean. To close, turn cap clockwise.



Fast acting relief
Lasts up to 12 hours, through
the day or night
Non-drowsy



MUCINEX SINUS MAX CLEAR AND COOL SEVERE CONGESTION

oxymetazoline hci spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:72854-218

Route of Administration NASAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY)

(OXYMETAZOLINE - UNII:8VLN5B44ZY)

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Inactive Ingredients

Ingredient Name Strength

CETALKONIUM CHLORIDE (UNII: 8547401N9D)

EDETATE DISODIUM (UNII: 7FLD91C86K)	
EUCALYPTOL (UNII: RV6J6604TK)	
LINOLEIC ACID (UNII: 9KJL21T0QJ)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
MYRISTIC ACID (UNII: 013V7S25AW)	
OLEIC ACID (UNII: 2UMI9U37CP)	
WATER (UNII: 059QF0KO0R)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
MYRISTALKONIUM CHLORIDE (UNII: 0W2550L75T)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BENZODODECINIUM CHLORIDE (UNII: Y5A751G47H)	
CAMPHOR, (-)- (UNII: 213N3S8275)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
GLYCINE (UNII: TE7660XO1C)	
PALMITIC ACID (UNII: 2V16EO95H1)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
LINOLENIC ACID (UNII: 0RBV727H71)	
PALMITOLEIC ACID (UNII: 209B6YPZ4I)	
SODIUM CARBONATE (UNII: 45P3261C7T)	

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:72854- 218-11	1 in 1 CARTON	06/01/2025			
1		22 mL in 1 BOTTLE, PUMP; 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	06/01/2025				

Labeler - RB Health (US) LLC (081049410)

Revised: 2/2025 RB Health (US) LLC