MUCINEX SINUS MAX SINUS PRESSURE SEVERE CONGESTION- oxymetazoline hci spray RB Health (US) LLC

Mucinex Sinus Max Sinus Pressure 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

Purpose:

Nasal Decongestant

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.

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

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.

Other information

■ store at 20-25°C (68-77°F)

Inactive ingredients

benzalkonium chloride, benzododecinium chloride, cetalkonium chloride, dibasic sodium phosphate anhydrous, edetate disodium, glycerin, monobasic sodium phosphate anhydrous, myristalkonium chloride, phosphorous pentoxide, propylene glycol, purified water, sodium oxide, sorbitol

Ouestions?

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

Dist. by: RB Health (US)



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Drug Facts (continued)

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

KEEP CARTON FOR FULL INFORMATION Please visit our website www.mucinex.com

s: www.reckitt.com/patents





Dist. by: RB Health (US) Parsippany NJ 07054-0224 @2025 RB Health 122424 Made in India

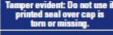
NDC 72854-219-11 SEVERE CONGESTION Oxymetazeline IICI 0.85% >> Masal Decongestant Nasal Spray /ORKING

Lasts 12 Hours

Powerful Sinus

Pressure Relief

3/4 FL OZ (22 mL)



How to Use



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

Prime Remove cap from tip. Depress several times until mist





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

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 SINUS PRESSURE SEVERE CONGESTION

oxymetazoline hci spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:72854-219

Route of Administration NASAL

Active Ingredient/Active Moiety

Basis of Strength Ingredient Name Strenath

OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY)	OXYMETAZOLINE	0.05 g
(OXYMETAZ OLINE - UNII:8VLN5B44ZY)	HYDROCHLORIDE	in 100 mL

Inactive Ingredients			
Ingredient Name	Strength		
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)			
BENZODODECINIUM CHLORIDE (UNII: Y5A751G47H)			
PHOSPHORUS PENTOXIDE (UNII: 51SWB7223J)			
CETALKONIUM CHLORIDE (UNII: 8547401N9D)			
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)			
GLYCERIN (UNII: PDC6A3C0OX)			
MYRISTALKONIUM CHLORIDE (UNII: 0W2550L75T)			
SODIUM OXIDE (UNII: 3075U8R23D)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
SORBITOL (UNII: 506T60A25R)			
EDETATE DISODIUM (UNII: 7FLD91C86K)			
SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM (UNII: 3980JIH2SW)			
WATER (UNII: 059QF0KO0R)			

ı	Packaging				
4	tem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:72854- 219-11	1 in 1 CARTON	06/01/2025		
]	L	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