MUCINEX NIGHTSHIFT SINUS MAXIMUM STRENGTH- acetaminophen, dextromethorphan hydrobromide, and triprolidine hydrochloride tablet, coated
RB Health (US) LLC

Mucinex Rapid+Clear Nighttime Headache, Cough & Fever

Active ingredients
(in each caplet)

Acetaminophen 325 mg
Dextromethorphan HBr 10 mg
Triprolidine HCl 1.25 mg

Uses
■ temporarily relieves these common cold and flu symptoms:
■ cough ■ minor aches and pains
■ sore throat ■ headache
■ runny nose ■ sneezing
■ itching of the nose or throat
■ itchy, watery eyes due to hay fever
■ temporarily reduces fever
■ controls cough to help you get to sleep

Warnings
Liver warning: This product contains acetaminophen.
Severe liver damage may occur if you take:
■ more than 8 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.
■ 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 ■ glaucoma
■ trouble urinating due to an enlarged prostate gland
■ a breathing problem such as emphysema or chronic bronchitis
■ persistent or chronic cough such as occurs with smoking, asthma, 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
■ taking sedatives or tranquilizers

When using this product

■ do not use more than directed
■ excitability may occur, especially in children
■ marked drowsiness may occur
■ alcohol, sedatives, and tranquilizers may increase drowsiness
■ avoid alcoholic drinks
■ use caution when driving amotor vehicle or operatingmachinery

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.

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 8 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)

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

Inactive ingredients
croscarmellose sodium, crospovidone, hypromellose, microcrystalline cellulose, polyethylene glycol, polysorbate 80, povidone, titanium dioxide

Acetaminophen 325 mg ..... Pain reliever/fever reducer
Dextromethorphan HBr 10 mg.....Cough suppressant
Triprolidine HCl 1.25 mg.....Antihistamine

Keep out of reach of children.

MUCINEX NIGHTSHIFT SINUS MAXIMUM STRENGTH

acetaminophen, dextromethorphan hydrobromide, and triprolidine hydrochloride tablet, coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72854-209
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH)	DEXTROMETHORPHAN	10 mg

(DEXTROMETHORPHAN - UNII:7355X3ROTS)	HYDROBROMIDE	10 mg
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
TRIPROLIDINE HYDROCHLORIDE (UNII: YAN7R5L890) (TRIPROLIDINE - UNII:2L8T9S52QM)	TRIPROLIDINE HYDROCHLORIDE	1.25 mg

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
CROSPOVIDONE, UNSPECIFIED (UNII: 2S7830E561)	

Product Characteristics

Color	yellow	**Score**	no score
Shape	OVAL	**Size**	19mm
Flavor		**Imprint Code**	VVV;LOGOcrescentmoonplus
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72854-209-20	2 in 1 CARTON	05/01/2025	
1		10 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	

Labeler - RB Health (US) LLC (081049410)

Revised: 5/2025 RB Health (US) LLC