MUCINEX DAY TIME PAIN, HEADACHE AND MUCUS CONGESTION AND MUCINEX NIGHT TIME HEADACHE, COUGH AND FEVER MAXIMUM STRENGTH-acetaminophen, dextromethorphan hydrobromide, guaifenesin, triprolidine hydrochloride
RB Health (US) LLC

MUCINEX® Rapid Clear® Pain, Headache & Mucus Congestion and Nighttime Caplets

Drug Facts

Acetaminophen 325 mg
Guaifenesin 200 mg

Mucinex Nightshift Night Sinus Purposes
Acetaminophen 325 mg Pain reliever/fever reducer
Dextromethorphan HBr 10 mg Cough suppressant
Triprolidine HCl 1.25 mg Antihistamine

Active ingredients (in each caplet)	***Purposes***
Mucinex Sinus-Max Day Pressure, Pain & Cough	
Acetaminophen 325 mg	**Pain reliever**
Guaifenesin 200 mg	Expectorant

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Dextromethorphan HBr 10 mg	Cough suppressant
Triprolidine HCl 1.25 mg	Antihistamine

Uses

Mucinex Sinus-Max Day Pressure, Pain & Cough

- temporarily relieves:
  - nasal congestion
  - headache
  - minor aches and pains
  - cough
  - sinus congestion and pressure
- temporarily promotes nasal and/or sinus drainage

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

Mucinex Nightshift Night Sinus

- temporarily relieves these common cold and flu symptoms:
  - cough
  - nasal congestion
  - minor aches and pains
  - sore throat
  - headache
  - sneezing
  - sinus congestion and pressure
  - runny nose
  - 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 4,000 mg in 24 hours, which is the maximum daily amount
- 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 (Nightshift Night Sinus only)

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
- heart disease
- diabetes
- high blood pressure
- thyroid disease
- glaucoma ***(Nightshift Night Sinus only)***
- trouble urinating due to an enlarged prostate gland
- a breathing problem such as emphysema or chronic bronchitis ***(Nightshift Night Sinus only)***
- 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
- taking sedatives or tranquilizers ***(Nightshift Night Sinus only)***

When using this product

- **do not use more than directed**
- excitability may occur, especially in children ***(Nightshift Night Sinus only)***
- marked drowsiness may occur ***(Nightshift Night Sinus only)***
- alcohol, sedatives, and tranquilizers may increase drowsiness ***(Nightshift Night Sinus only)***
- avoid alcoholic drinks ***(Nightshift Night Sinus only)***
- use caution when driving a motor vehicle or operating machinery ***(Nightshift Night Sinus only)***

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- pain, nasal congestion, 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 fever, rash, or headache that lasts. These could be signs of a serious condition.

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

Keep out of reach of children.

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

Mucinex Sinus-Max Day Pressure, Pain & Cough

- **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

Mucinex Nightshift Night Sinus

- **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)

Inactive ingredients

Mucinex Sinus-Max Day Pressure, Pain & Cough

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

Mucinex Nightshift Night Sinus

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

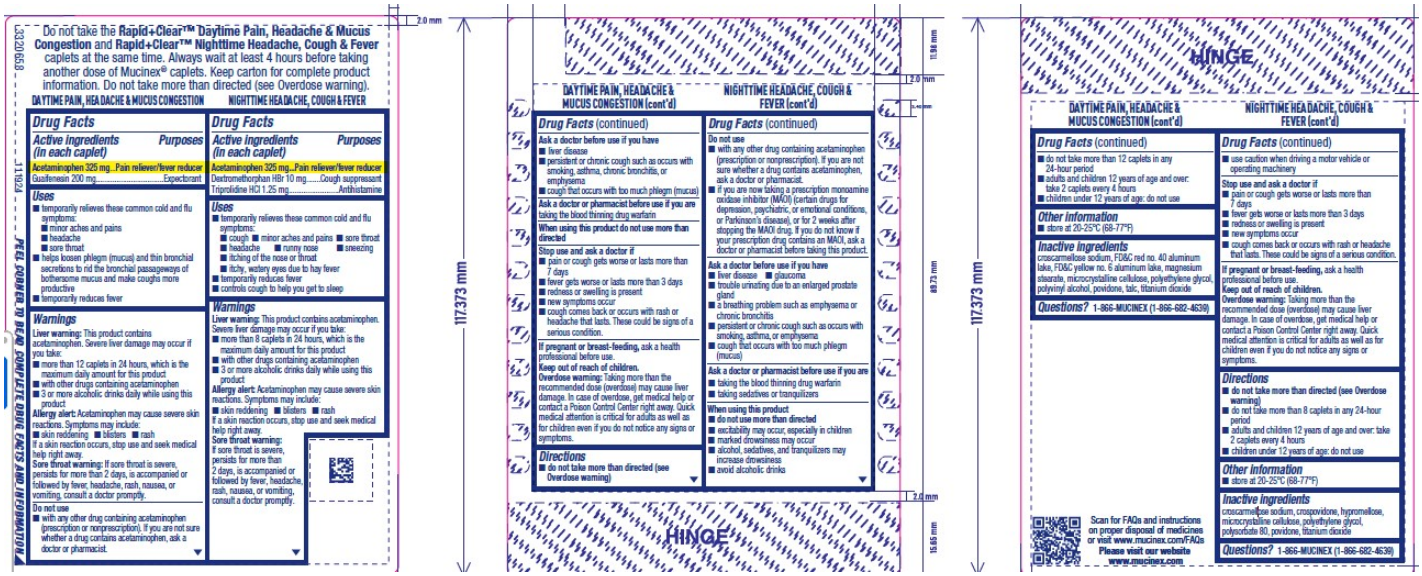
Questions? 1-866-MUCINEX (1-866-682-4639)

You may also report side effects to this phone number.

Dist. by: RB Health (US)
Parsippany, NJ 07054-0224

PRINCIPAL DISPLAY PANEL - Kit Carton

NDC 72854-210



MUCINEX DAY TIME PAIN, HEADACHE AND MUCUS CONGESTION

AND MUCINEX NIGHT TIME HEADACHE, COUGH AND FEVER MAXIMUM STRENGTH

acetaminophen, dextromethorphan hydrobromide, guaifenesin, triprolidine hydrochloride kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72854-210

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72854-210-10	1 in 1 CARTON	07/01/2025	
1		1 in 1 KIT		
2	NDC:72854-210-20	2 in 1 CARTON	07/01/2025	
2		1 in 1 KIT		
3	NDC:72854-210-40	4 in 1 CARTON	07/01/2025	
3		1 in 1 KIT		

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	1 BLISTER PACK	6
Part 2	1 BLISTER PACK	4

Part 1 of 2

MAXIMUM STRENGTH MUCINEX SINUS-MAX PRESSURE, PAIN AND COUGH

acetaminophen, guaifenesin tablet, film coated

Product Information	
Item Code (Source)	NDC:72854-208
Route of Administration	ORAL

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

Inactive Ingredients	
Ingredient Name	**Strength**
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

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		6 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	07/01/2025	

Part 2 of 2

MUCINEX NIGHTSHIFT SINUS MAXIMUM STRENGTH

acetaminophen, dextromethorphan hydrobromide, and triprolidine hydrochloride tablet, coated

Product Information

Item Code (Source)	NDC:72854-209
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
TRIPROLIDINE HYDROCHLORIDE (UNII: YAN7R5L890) (TRIPROLIDINE -	TRIPROLIDINE	1.25 mg

UNII:2L8T9S52QM)	HYDROCHLORIDE	1.25 mg

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

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		4 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	07/01/2025	

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: 6/2025 RB Health (US) LLC