

**MUCINEX RAPID CLEAR PAIN, HEADACHE AND MUCUS CONGESTION AND NIGHTTIME LIQUID GELS- acetaminophen, guaifenesin, dextromethorphan hydrobromide, doxylamine succinate
RB Health (US) LLC**

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

Day:

Acetaminophen 325 mg

Guaifenesin 200 mg

Night:

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Doxylamine succinate 6.25 mg

Day:

Acetaminophen 325 mg...Pain reliever/fever reducer

Guaifenesin 200 mg.....Expectorant

Night:

Acetaminophen 325 mg...Pain reliever/fever reducer

Dextromethorphan HBr 10 mg.....Cough suppressant

Doxylamine succinate 6.25 mg..... Antihistamine

Day:

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

Night:

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

Day:

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.

Ask a doctor before use if you have

- liver disease
- 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

When using this product do not use more than directed

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.

Night:

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.
- 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 a motor vehicle or operating machinery

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.

Day:

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

Night:

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

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.

Other information

■ store at 20-25°C (68-77°F)

■ avoid excessive heat

Day:

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

Night:

Inactive ingredients

D&C yellow no. 10, FD&C blue no. 1, gelatin, glycerin, hypromellose, isopropyl alcohol, lecithin (sunflower), light mineral oil, mannitol, polyethylene glycol 400, povidone, propylene glycol, purified water, 1,4-sorbitan, sorbitol, titanium dioxide

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

MUCINEX RAPID CLEAR PAIN, HEADACHE AND MUCUS CONGESTION AND NIGHTTIME LIQUID GELS

acetaminophen, guaifenesin, dextromethorphan hydrobromide, doxylamine succinate kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72854-217
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72854-217-24	2 in 1 CARTON	06/01/2025	
1		1 in 1 BLISTER PACK; Type 9: Other Type of Part 3 Combination Product (e.g., Drug/Device/Biological Product)		

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BLISTER PACK	16
Part 2	1 BLISTER PACK	8

Part 1 of 2

MUCINEX RAPID CLEAR DAYTIME PAIN, HEADACHE AND MUCUS CONGESTION

acetaminophen, guaifenesin capsule, liquid filled

Product Information

Item Code (Source)	NDC:72854-001
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Route of Administration	ORAL
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Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POVIDONE (UNII: FZ989GH94E)	
GELATIN (UNII: 2G86QN327L)	

GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	
LECITHIN, SUNFLOWER (UNII: 834K0WOS5G)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Product Characteristics

Color	blue	Score	no score
Shape	OVAL	Size	23mm
Flavor		Imprint Code	AR09
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		16 in 1 BLISTER PACK; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012		

Part 2 of 2

MUCINEX RAPID CLEAR NIGHTTIME HEADACHE, COUGH AND FEVER

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate capsule, liquid filled

Product Information

Item Code (Source)	NDC:72854-002
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	10 mg

(DEXTROMETHORPHAN - UNII:7355X3ROTS)	HYDROBROMIDE	10 mg
DOXYLAMINE SUCCINATE (UNII: V9B19B5Y12) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg

Inactive Ingredients

Ingredient Name	Strength
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
1,4-SORBITAN (UNII: AV0YTZ4E6J)	
LECITHIN, SUNFLOWER (UNII: 834K0WOS5G)	
WATER (UNII: 059QF0K00R)	
SORBITOL (UNII: 506T60A25R)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
GELATIN (UNII: 2G86QN327L)	
MANNITOL (UNII: 3OWL53L36A)	
POVIDONE (UNII: FZ989GH94E)	

Product Characteristics

Color	green	Score	no score
Shape	OVAL	Size	23mm
Flavor		Imprint Code	AR17
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		8 in 1 BLISTER PACK; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012		

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: 11/2025

RB Health (US) LLC