MAXIMUM STRENGTH MUCINEX FAST-MAX DAY TIME COLD AND FLU AND NIGHT TIME COLD AND FLU- acetaminophen, dextromethorphan hydrobromide, diphenhydramine hydrochloride, guaifenesin, and phenylephrine hydrochloride RB Health (US) LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Maximum Strength Mucinex® Fast-Max® Day Time Cold & Flu and Night Time Cold & Flu Drug Facts

Active ingredients (in each caplet) Mucinex FAST-MAX DAY TIME COLD & FLU	Purposes
Acetaminophen 325 mg	Pain reliever/fever reducer
Dextromethorphan HBr 10mg	Cough suppressant
Guaifenesin 200 mg	Expectorant
Phenylephrine HCl 5 mg	Nasal decongestant
Active ingredients (in each caplet)	Purposes
Mucinex FAST-MAX NIGHT TIME COLD & FLU	
Acetaminophen 325 mg	Pain reliever/fever reducer
Diphenhydramine HCl 12.5	Antihistamine/cough
mg Phenylephrine HCl 5 mg	suppressant Nasal decongestant

Uses

Mucinex Fast-Max DAY TIME COLD & FLU

- temporarily relieves these common cold and flu symptoms:
 - cough
 - minor aches and pains
 - headache
 - nasal congestion
 - sore throat
 - sinus congestion and pressure
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- controls cough to help you get to sleep
- temporarily reduces fever

Mucinex Fast-Max NIGHT TIME COLD & FLU

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

Warnings

Liver warning

This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 12 caplets 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

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.
- with any other product containing diphenhydramine, even one used on the skin (NIGHT TIME only)
- 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
- trouble urinating due to an enlarged prostate gland

- glaucoma (NIGHT TIME only)
- a breathing problem such as emphysema or chronic bronchitis (*NIGHT TIME 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 (NIGHT TIME only)

When using this product

- do not use more than directed
- excitability may occur, especially in children (NIGHT TIME only)
- marked drowsiness may occur (NIGHT TIME only)
- alcohol, sedatives, and tranquilizers may increase drowsiness (NIGHT TIME only)
- avoid alcoholic drinks (NIGHT TIME only)
- be careful when driving a motor vehicle or operating machinery (NIGHT TIME 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

- 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

Other information

■ store at 20-25°C (68-77°F)

Inactive ingredients

(Mucinex FAST-MAX DAY TIME COLD & FLU)

croscarmellose sodium, crospovidone, 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

Inactive ingredients

(Mucinex FAST-MAX NIGHT TIME COLD & FLU)

croscarmellose sodium, crospovidone, FD&C blue no. 1 aluminum lake, FD&C blue no. 2 aluminum lake, ferric oxide yellow, methacrylic acid – ethyl acrylate copolymer (1:1) type A, mica, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, sodium bicarbonate, talc, titanium dioxide

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

PRINCIPAL DISPLAY PANEL - Kit Carton

NDC 63824-511-40

MAXIMUM STRENGTH

Mucinex®

FAST-MAX®

DAY

TIME

COLD & FLU

Acetaminophen – Pain Reliever/Fever Reducer

 $Dextromethor phan\ HBr-Cough\ Suppressant$

Guaifenesin – Expectorant

Phenylephrine HCl – Nasal Decongestant

HEADACHE

BODY PAIN

SORE THROAT

FEVER

CHEST CONGESTION

COUGH

ALL IN

ONE*

NASAL CONGESTION

SINUS CONGESTION

SINUS PRESSURE

Actual Size

FOR AGES 12+

24 CAPLETS

NIGHT

TIME

COLD & FLU

Acetaminophen – Pain Reliever/Fever Reducer

Diphenhydramine HCl – Antihistamine/

Cough Suppressant

Phenylephrine HCl – Nasal Decongestant

HEADACHE

BODY PAIN

SORE THROAT

FEVER

ITCHY THROAT COUGH ALL IN ONE* NASAL CONGESTION SNEEZING RUNNY NOSE

Actual Size

FOR AGES 12+

16 CAPLETS

TOTAL 40 CAPLETS



Tamper evident: Do not use if carton is damaged or if printed seal on blister is broken or missing.

NDC 63824-511-40

COLD & FLU FAST-MAX O NIGHT TIME

Maximum Strength per 4-hour dose Do not take more than a total of 12 captes in any 24-hour period "DAY TIME COLD & FLU helps to releve these symptoms day or night NIGHT TIME COLD & FLU helps to relieve these symptoms at night

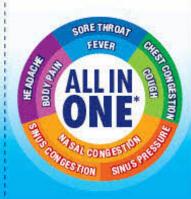
Take only as directed. Keep carton for full information.





COLD & FLU

Acetaminophen – Pain Reliever/Fe Dextromethorphan HBr – Cough Su Guardenesin – Expectorant



Aotual Size

FOR AGES 12+



TOTAL 40 CAPLETS

HEADACHE WEAL CONGESTION RUNIN HOS

COLD & FLU

Acetaminophen – Pain Reliever/Fever Reducer Diphenhydramine HCl – Antihistamine/

Phenylephrine HCI - Nasal Decongestant

SORE THROAT

FEVER

Aotual Size



FOR AGES 12+



Questions? 1-866-Muchubi (1-866-882-4839) You may aso report ade effects to this phone number

www.mucinex.com Patents: www.rb.com/patents

Dist. by: RB Heath (US) Parsippany, NJ 07054-0224 C2020 RB Heath

35/569 08020

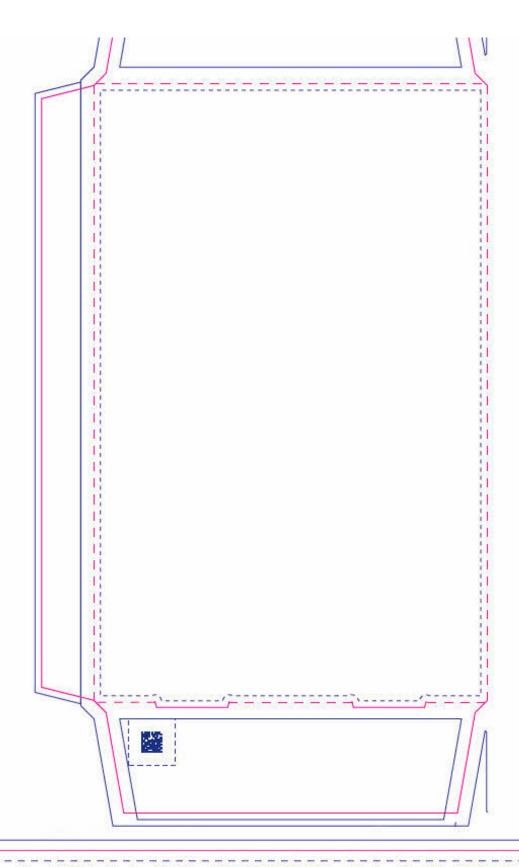
M 9951 63824

CAPLETS

LOT:

EXP: MADE IN:

3157569



313845

Do not take the Mucinex Fast-Max DAY TIME COLD & FLU and Mucinex Fast-Max NIGHT TIME COLD & FLU caplets at the same time. Always wait at least 4 hours before taking another dose of Mucinex caplets. Do not take more than a total of 12 caplets in any 24-hour period.

Take only as directed.

Drug Facts

Active ingredients (in each caplet)

Purposes

Reseal Area

Drug Facts (continued)

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.
- with any other product containing diphenhydramine, even one used on the skin (NIGHT TIME

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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
- trouble urinating due to an enlarged prostate gland
- glaucoma (NIGHT TIME only)
- a breathing problem such as emphysema or chronic bronchitis (NIGHT TIME 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 (NIGHT TIME only)

When using this product

- do not use more than directed
- mexcitability may occur, especially in children (NIGHT TIME only)
- marked drowsiness may occur (NIGHT TIME only)
- alcohol, sedatives, and tranquilizers may increase drowsiness (NIGHT TIME only)
- avoid alcoholic drinks (NIGHT TIME only)
- be careful when driving a motor vehicle or operating machinery (NIGHT TIME 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.

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Drug Facts (continued)

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 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

Other information

■ store at 20-25°C (68-77°F)

Inactive ingredients

(Mucinex FAST-MAX DAY TIME COLD & FLU)

croscarmellose sodium, crospovidone, 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

Inactive ingredients

(Mucinex FAST-MAX NIGHT TIME COLD & FLU)

croscarmellose sodium, crospovidone, FD&C blue no. 1 aluminum lake, FD&C blue no. 2 aluminum lake, ferric oxide yellow, methacrylic acid — ethyl acrylate copolymer (1:1) type A, mica, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, sodium bicarbonate, talc, titanium dioxide

Questions?

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

You may also report side effects to this phone number.



MAXIMUM STRENGTH MUCINEX FAST-MAX DAY TIME COLD AND FLU AND NIGHT TIME COLD AND FLU

acetaminophen, dextromethorphan hydrobromide, diphenhydramine hydrochloride, guaifenesin, and phenylephrine hydrochloride kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:63824-511

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:63824-511-10	1 in 1 CARTON	06/01/2020		
2	NDC:63824-511-20	2 in 1 CARTON	06/01/2020		
2		1 in 1 KIT			
3	NDC:63824-511-40	4 in 1 CARTON	06/01/2020		
3		1 in 1 KIT			

Quant	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 FAST-MAX DAY TIME COLD AND FLU

acetaminophen, dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride tablet

Product Information Route of Administration ORAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
Acetaminophen (UNII: 36209 ITL9D) (Acetaminophen - UNII: 36209 ITL9D)	Acetaminophen	325 mg		
Guaifenesin (UNII: 495W7451VQ) (Guaifenesin - UNII:495W7451VQ)	Guaifenesin	200 mg		
Dextromethorphan Hydrobromide (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	Dextro metho rphan Hydro bro mide	10 mg		
Phenylephrine Hydrochloride (UNII: 04JA59TNSJ) (Phenylephrine - UNII:1WS297W6MV)	Phenylephrine Hydrochloride	5 mg		

Inactive Ingredients	
Ingredient Name	Strength
croscarmellose sodium (UNII: M28OL1HH48)	
CROSPOVIDONE (UNII: 2S7830E561)	
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: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
PO VIDO NE, UNSPECIFIED (UNII: FZ989GH94E)	
talc (UNII: 7SEV7J4R1U)	
titanium dioxide (UNII: 15FIX9V2JP)	

Product Characteristics				
Color RED Score no score				
Shape	OVAL	Size	20 mm	
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	Marketing Start Date	Marketing End Date		
OTC monograph final	part341	06/01/2020		

Part 2 of 2

MAXIMUM STRENGTH MUCINEX FAST-MAX NIGHT TIME COLD AND FLU

acetaminophen, diphenhydramine hydrochloride, and phenylephrine hydrochloride tablet

Product Information

Route of Administration ORAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
Acetaminophen (UNII: 36209ITL9D) (Acetaminophen - UNII:36209ITL9D)	Acetaminophen	325 mg		
Diphenhydramine Hydrochloride (UNII: TC2D6JAD40) (Diphenhydramine - UNII:8GTS82S83M)	Diphenhydramine Hydrochloride	12.5 mg		
Phenylephrine Hydrochloride (UNII: 04JA59TNSJ) (Phenylephrine - UNII:1WS297W6MV)	Phenylephrine Hydrochloride	5 mg		

Inactive Ingredients	
Ingredient Name	Strength
croscarmellose sodium (UNII: M28 OL 1HH48)	
CROSPOVIDONE (UNII: 2S7830E561)	
FD&C blue no. 1 (UNII: H3R47K3TBD)	
FD&C blue no. 2 (UNII: L06K8R7DQK)	
ALUMINUM O XIDE (UNII: LMI26 O 69 33)	
ferric oxide yellow (UNII: EX438O2MRT)	
METHACRYLIC ACID - ETHYL ACRYLATE COPOLYMER (1:1) TYPE A (UNII: NX76LV5T8J)	
mica (UNII: V8A1AW0880)	

microcrystalline cellulose (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
PO VIDO NE, UNSPECIFIED (UNII: FZ989 GH94E)	
sodium bicarbonate (UNII: 8MDF5V39QO)	
talc (UNII: 7SEV7J4R1U)	
titanium dioxide (UNII: 15FIX9 V2JP)	

Product Characteristics				
Color	BLUE	Score	no score	
Shape	OVAL	Size	20 mm	
Flavor		Imprint Code	VVV;CF	
Contains				

	Packaging							
Ш	# 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 final	part341	06/01/2020				

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part341	06/01/2020				

Labeler - RB Health (US) LLC (081049410)

Revised: 10/2020 RB Health (US) LLC