MUCINEX FAST-MAX LIQUID GELS DAY NIGHT COLD AND FLU- acetaminophen, dextromethorphan hydrobromide, and doxylamine succinate
RB Health (US) LLC

MUCINEX® FAST-MAX® Liquid Gels - Day Night Cold & Flu

Purpose Day:
Acetaminophen 325 mg...Pain reliever/fever reducer

Dextromethorphan HBr 10 mg.....Cough suppressant

Purpose Night:

Acetaminophen 325 mg...Pain reliever/fever reducer
Dextromethorphan HBr 10 mg.....Cough suppressant
Doxylamine succinate 6.25 mg..... Antihistamine

Day:

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Night:

Acetaminophen 325 mg
Dextromethorphan HBr 10 mg
Doxylamine succinate 6.25 mg

Day:

■ temporarily relieves these common cold and flu symptoms:
■ cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
■ the intensity of coughing
■ the impulse to cough to help you get to sleep
■ minor aches and pains ■ sore throat
■ headache
■ temporarily reduces fever

Night:

■ 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.

■ 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
■ 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

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.

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.

Night:

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

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

Other information
■ store at 20-25°C (68-77°F)
■ avoid excessive heat

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.

Day:

FD&C red no. 40, FD&C yellow no. 6, 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

Night:

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)

FAST DISSOLVING
LIQUID GELS

NEW FORMULAS | MAXIMUM STRENGTH | NEW FORMULAS | NEW FORMULAS

NDC 72854-168-24

Mucinex®

FAST-MAX®

DAY TIME COLD & FLU

Acetaminophen » Pain Reliever/Fever Reducer
Dextromethorphan HBr » Cough Suppressant

NIGHT TIME COLD & FLU

Acetaminophen » Pain Reliever/Fever Reducer
Dextromethorphan HBr » Cough Suppressant
Doxylamine succinate » Antihistamine

AGES 12+

AGES 12+

ACTUAL SIZE

16 LIQUID GELS (Liquid Filled Capsules)

TOTAL 24 LIQUID GELS

ACTUAL SIZE

8 LIQUID GELS (Liquid Filled Capsules)

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

PARENTS: Learn about teen medicine abuse www.StopMedicineAbuse.org

how2recycle.info

MULTI-LAYER TRAY

PAPER BOX

3 63824 99902 9

MAXIMUM STRENGTH
Mucinex FAST-MAX
DAY TIME COLD & FLU
NIGHT TIME COLD & FLU

*Fast-Max® Daytime Cold & Flu helps to relieve these symptoms day or night; Fast-Max® Nighttime Cold & Flu helps to relieve these symptoms at night

Maximum Strength per 4-hour dose

Take only as directed.

Keep carton for full information.

reckitt

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

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Patents: www.reckitt.com/patents

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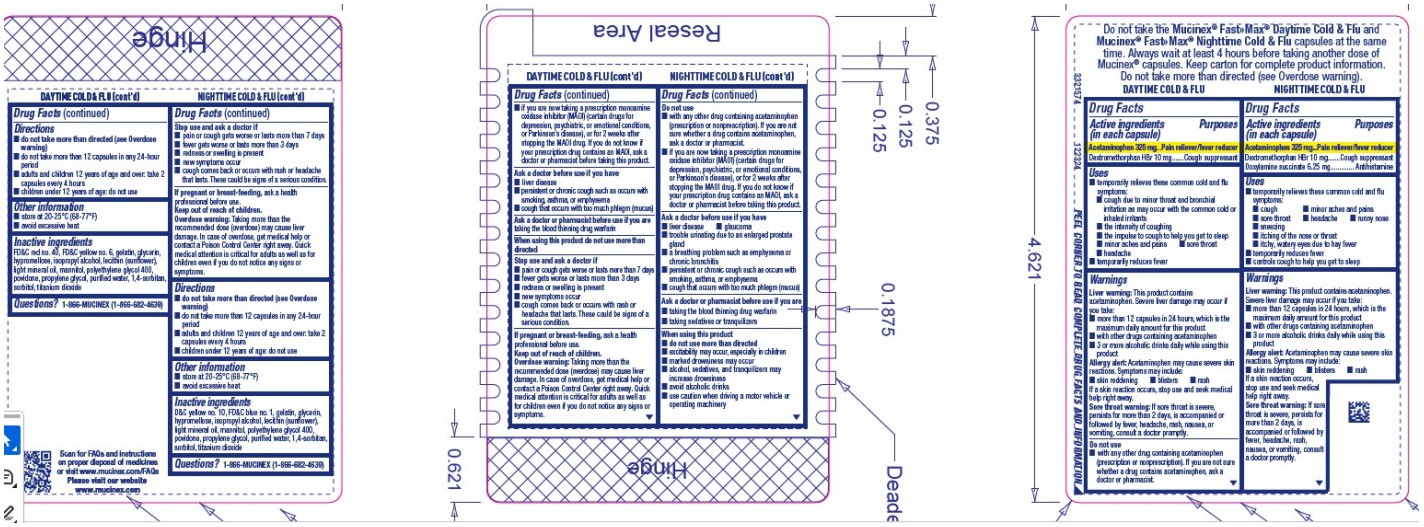

MUCINEX FAST-MAX LIQUID GELS DAY NIGHT COLD AND FLU

acetaminophen, dextromethorphan hydrobromide, and doxylamine succinate kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72854-168

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72854-168-10	1 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)		
2	NDC:72854-168-24	2 in 1 CARTON	06/01/2025	
2		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	0 BLISTER PACK	1
Part 2	0 BLISTER PACK	1

Part 1 of 2

MUCINEX FAST-MAX COLD AND FLU MAXIMUM STRENGTH

acetaminophen, dextromethorphan hydrobromide capsule, liquid filled

Product Information

Item Code (Source)	NDC:72854-171

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

Inactive Ingredients

Ingredient Name	Strength
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN (UNII: 2G86QN327L)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
MANNITOL (UNII: 3OWL53L36A)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POVIDONE (UNII: FZ989GH94E)	
WATER (UNII: 059QF0KO0R)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SORBITOL (UNII: 506T60A25R)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
1,4-SORBITAN (UNII: AV0YTZ4E6J)	
LECITHIN, SUNFLOWER (UNII: 834K0WOS5G)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
GLYCERIN (UNII: PDC6A3C0OX)	

Product Characteristics

Color	red	**Score**	no score
Shape	OVAL	**Size**	24mm
Flavor		**Imprint Code**	AR18
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72854-171-08	1 in 1 CARTON		
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	06/01/2025	

Part 2 of 2

MUCINEX FAST-MAX NIGHTTIME COLD AND FLU

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate capsule, liquid filled

Product Information

Item Code (Source)	NDC:72854-003
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
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg

Inactive Ingredients

Ingredient Name	Strength
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
SORBITOL (UNII: 506T60A25R)	
1,4-SORBITAN (UNII: AV0YTZ4E6J)	
POVIDONE (UNII: FZ989GH94E)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
MANNITOL (UNII: 3OWL53L36A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
WATER (UNII: 059QF0KO0R)	
LECITHIN, SUNFLOWER (UNII: 834K0WOS5G)	

Product Characteristics

Color	green	**Score**	no score
Shape	OVAL	**Size**	24mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		1 in 1 CARTON		
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		

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