MUCINEX FAST-MAX COLD AND FLU- acetaminophen, dextromethorphan hydrobromide tablet
RB Health (US) LLC

Mucinex Fast-Max Cold & Flu

Drug Facts

Acetaminophen 325 mg...Pain reliever/fever reducer
Dextromethorphan HBr 10 mg...Cough suppressant

Uses

Uses
■ 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

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 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

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

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

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
croscarmellose sodium, FD&C red no. 40 aluminum lake, FD&C yellow no. 6 aluminum lake, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, talc, titanium dioxide

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

Keep out of reach of children.

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

MUCINEX FAST-MAX COLD AND FLU

acetaminophen, dextromethorphan hydrobromide tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72854-161
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 YELLOW NO. 6 ALUMINUM LAKE (UNII: GYP6Z2JR6Q)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FD&C RED NO. 40 ALUMINUM LAKE (UNII: 6T47AS764T)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
POVIDONE (UNII: FZ989GH94E)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
TALC (UNII: 7SEV7J4R1U)	

Product Characteristics

Color	red	**Score**	no score
Shape	OVAL	**Size**	8mm
Flavor		**Imprint Code**	Chevron
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72854-161-10	1 in 1 CARTON	04/01/2025	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:72854-161-20	2 in 1 CARTON	04/01/2025	
2		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:72854-161-02	2 in 1 POUCH; Type 0: Not a Combination Product	04/01/2025	
4	NDC:72854-161-04	2 in 1 CARTON	04/01/2025	
4		2 in 1 POUCH; 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	04/01/2025	

Labeler - RB Health (US) LLC (081049410)

Revised: 3/2025 RB Health (US) LLC