

MUCINEX FAST-MAX COLD AND FLU MAXIMUM STRENGTH- acetaminophen, dextromethorphan hydrobromide capsule, liquid filled
RB Health (US) LLC

Mucinex[®] Fast-Max[®] Cold & Flu

Maximum Strength

Drug Facts

Active ingredients (in each liquid gel) Purposes
Acetaminophen 325 mg Pain reliever/fever reducer
Dextromethorphan HBr 10 mg Cough suppressant

<i>Active ingredients (in each liquid gel)</i>	<i>Purposes</i>
Acetaminophen 325 mg	Pain reliever/fever reducer
Dextromethorphan HBr 10 mg	Cough suppressant

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

This product contains acetaminophen. Severe liver damage may occur if you take:
more than 12 liquid gels in 24 hours, which is the maximum daily amount
with other drugs containing acetaminophen
3 or more alcoholic drinks daily while using this product

Liver warning

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

- more than 12 liquid gels 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.
- 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
- 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.

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 liquid gels in any 24-hour period
- adults and children 12 years of age and over: take 2 liquid gels 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

Inactive ingredients

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

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 - 16 Capsule Blister Pack Carton

Fast Dissolving Liquid Gels!

NDC 72854-171

MAXIMUM STRENGTH

MUCINEX®

FAST-MAX®

COLD & FLU

Acetaminophen – Pain Reliever/Fever Reducer

Dextromethorphan HBr – Cough Suppressant

LIQUID GELS

(Liquid Filled Capsules)

Actual Size

FOR AGES 12 +

FAST DISSOLVING LIQUID GELS



NEW FORMULA

MAXIMUM STRENGTH

NEW FORMULA

NEW FORMULA

NDC 72854-171-16

Mucinex®

FAST»MAX®

COLD & FLU

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

MAXIMUM STRENGTH

FAST»MAX®

COLD & FLU

* Helps to relieve these symptoms
day or night

Maximum Strength per 4-hour
dose

Do not take more than a total of
12 capsules in any 24-hour period.
Take only as directed.

Keep carton for full information.



Scan for FAQs and instructions on
proper disposal of medicines or
visit www.mucinex.com/FAQs
Please visit our website
www.mucinex.com

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

how2recycle.info



PAPER BOX

MULTI-LAYER TRAY



3 63824 99900 5

ALL IN ONE*

FEVER

SORE THROAT

COUGH

MINOR PAINS

BODY ACHES

HEADACHE



16 LIQUID GELS
(Liquid Filled Capsules)

ACTUAL SIZE

AGES 12+

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(in each capsule)****Purposes**

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Drug Facts (continued)

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

**reckitt**

Patents:
www.reckitt.com/patents
Dist. by: RB Health (US)
Parsippany, NJ 07054-0224

121824
3321582

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MUCINEX FAST-MAX COLD AND FLU MAXIMUM STRENGTH

acetaminophen, dextromethorphan hydrobromide capsule, liquid filled

Product Information				
Product Type		HUMAN OTC DRUG	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
SORBITOL (UNII: 506T60A25R)				
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)				
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)				
GLYCERIN (UNII: PDC6A3C0OX)				
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)				
ISOPROPYL ALCOHOL (UNII: ND2M416302)				
LIGHT MINERAL OIL (UNII: N6K5787QVP)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
WATER (UNII: 059QF0KO0R)				
LECITHIN, SUNFLOWER (UNII: 834K0WOS5G)				
MANNITOL (UNII: 3OWL53L36A)				
1,4-SORBITAN (UNII: AV0YTZ4E6J)				
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:72854-171-16	2 in 1 CARTON	06/01/2025	
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:72854-171-08	1 in 1 CARTON	06/01/2025	
2		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
	NDC 72854			

3	NDC: 72854-171-10	1 in 1 CARTON	06/01/2025	
3		10 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		06/01/2025	

Labeler - RB Health (US) LLC (081049410)

Revised: 6/2025

RB Health (US) LLC