

**MUCUS RELIEF CONGESTION AND HEADACHE MAXIMUM STRENGTH-
acetaminophen, guaifenesin, phenylephrine hcl liquid
Dolgenercorp, Inc. (DOLLAR GENERAL & REXALL)**

Drug Facts

Active ingredients (in each 20 mL)

Acetaminophen 650 mg

Guaifenesin 400 mg

Phenylephrine HCL 10 mg

Purposes

Pain reliever/fever reducer

Expectorant

Nasal decongestant

Uses

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

Warnings

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

- more than 4,000 mg of acetaminophen in 24 hours
- 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.
- for children under 12 years of age
- 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
- thyroid disease
- high blood pressure
- 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

- 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 rash or persistent headache. 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 (1-800-222-1222) 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 6 doses in any 24-hour period
- measure only with dosing cup provided. Do not use any other dosing device.
- keep dosing cup with products
- mL = milliliter
- dose as follows or as directed by a doctor
- adults and children 12 years and older: 20 mL in dosing cup provided every 4 hours
- children under 12 years of age: do not use

Other information

- **each 20 mL contains:** sodium 12 mg
- store between 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

anhydrous citric acid, EDTA disodium, FD&C blue #1, FD&C red #40, flavor, glycerin, propyl gallate, propylene glycol, purified water, sodium benzoate, sorbitol, sucralose, trisodium citrate dihydrate, xanthan gum

Principal Display Panel

Compare to the active ingredients of Maximum Strength Mucinex® Fast-Max® Congestion & Headache*

Maximum Strength

Fast Acting

Mucus Relief

Congestion & Headache

Acetaminophen

Guaifenesin

Phenylephrine HCL

Pain Reliever/Fever Reducer

Expectorant

Nasal Decongestant

For ages 12 years and over

FL OZ (mL)

*This product is not manufactured or distributed by Reckitt Benckiser, distributor of Maximum Strength Mucinex® Fast-Max® Congestion & Headache.

**TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS
BROKEN OR MISSING.**

DISTRIBUTED BY OLD EAST MAIN CO.

100 MISSION RIDGE

GOODLETTSVILLE, TN 37072

Package Label

DG health

Compare to the active ingredients of Maximum Strength Mucinex® Fast-Max® Congestion & Headache*

Maximum Strength Fast Acting Mucus Relief Congestion & Headache

Acetaminophen
Guaifenesin
Phenylephrine HCl
Pain Reliever/Fever Reducer
Expectorant
Nasal Decongestant

For ages 12 years and over

DISTRIBUTED BY OLD EAST MAIN CO.
100 MISSION RIDGE
GOODLETTSVILLE, TN 37072

100* Satisfaction Guaranteed! (888) 309-9030

A0513 PLD-B281D LB007843



9 fl oz
(266 mL)

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PLD-C281D
LB005829

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PEEL CORNER FOR MORE DRUG FACTS

Drug Facts (continued)

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Drug Facts (continued)

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DOLLAR GENERAL Mucus Relief Maximum Strength Congestion & Headache

MUCUS RELIEF CONGESTION AND HEADACHE MAXIMUM STRENGTH

acetaminophen, guaifenesin, phenylephrine hcl liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-757
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg in 20 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 20 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYL GALLATE (UNII: 8D45NN7V92)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-757-09	266 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	06/01/2019	

Labeler - Dolgencorp, Inc. (DOLLAR GENERAL & REXALL) (068331990)

Revised: 3/2024

Dolgencorp, Inc. (DOLLAR GENERAL & REXALL)