

**MUCUS RELIEF CONGESTION AND COUGH MAXIMUM STRENGTH-
dextromethorphan hbr, guaifenesin, phenylephrine hci liquid
TARGET Corporation**

Drug Facts

Active ingredients (in each 20 mL)

Dextromethorphan HBr 20 mg

Guaifenesin 400 mg

Phenylephrine HCl 10 mg

Purposes

Cough suppressant

Expectorant

Nasal decongestant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- temporarily relieves
 - cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritant
 - the intensity of coughing
 - the impulse to cough to help you get to sleep
 - nasal congestion due to a cold

Warnings

Do not use

- if you are 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.
- for children under 12 years of age

Ask a doctor before use if you have

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland

- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, emphysema
- cough that occurs with too much phlegm (mucus)

When using this product

do not use more than directed.

Stop use and ask a doctor if

- nervousness, dizziness or sleeplessness occur
- symptoms do not get better within 7 days or occur with fever
- 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.

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

- 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 product
- mL= milliliter
- shake well before using
- adult and children 12 years of age 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 17 mg**
- store between 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

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

Questions or comments?

Call 1-800-910-6874

Principal Display Panel

Compare to active ingredients in Maximum Strength Mucinex® Fast-Max® Severe Congestion & Cough*

maximum strength

fast mucus relief severe congestion and cough

dextromethorphan HBr 20 mg (cough suppressant)

guaifenesin 400 mg (expectorant)

phenylephrine HCl 10 mg (nasal decongestant)

controls cough

relieves nasal and chest congestion

thins and loosens mucus

AGES 12 + YEARS

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

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

Dist. by Target Corp.

Minneapolis, MN 55403

Product Label

NDC 11673-337-06

Compare to active ingredients in Maximum Strength Mucinex® Fast-Max® Severe Congestion & Cough*

maximum strength
fast mucus relief
severe congestion
and cough

dextromethorphan HBr 20 mg
(cough suppressant)
guaifenesin 400 mg (expectorant)
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(nasal decongestant)

controls cough
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AGES
12+
YEARS

6 FL OZ (177 mL)

PLD-B283D LB004222

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

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PEEL CORNER FOR MORE DRUG FACTS

Drug Facts (continued)

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Drug Facts (continued)

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PEEL CORNER FOR MORE DRUG FACTS

TARGET Maximum Strength Mucus Relief Severe Congestion and Cough

MUCUS RELIEF CONGESTION AND COUGH MAXIMUM STRENGTH

dextromethorphan hbr, guaifenesin, phenylephrine hci liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-337
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 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)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-337-06	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/31/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/31/2016	

