UP AND UP MAXIMUM STRENGTH FAST MUCUS RELIEF SEVERE CONGESTION AND COUGH- dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride liquid
Target Corporation
Reference Label Set Id: 7f4ccd7c-9b17-428b-8cdf-42237728f9b2

Up and Up maximum strength fast mucus relief severe congestion and cough

Drug Facts

Active ingredients (in each 20 mL)	***Purposes***
Dextromethorphan HBr 20 mg	Cough suppressant
Guaifenesin 400 mg	Expectorant
Phenylephrine HCl 10 mg	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 irritants
  - the intensity of coughing
  - the impulse to cough to help you get to sleep
  - nasal congestion due to a cold

Warnings

Do not use

- 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

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

Directions

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided
- do not use dosing cup with other products
- dose as follows or as directed by a doctor
- mL = milliliter
- **Adults 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 8 mg**
- store between 20-25°C (68-77°F)
- do not refrigerate
- dosing cup provided

Inactive ingredients

anhydrous citric acid, edetate disodium, FD&C Blue #1, FD&C Red #40, flavors, potassium citrate, propylene glycol, propyl gallate, purified water, sodium benzoate, sorbitol, sucralose, xanthan gum

Questions or comments?

(1-800-910-6874)

PRINCIPAL DISPLAY PANEL

NDC 11673-738-06

Compare to active ingredients in **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

6 FL OZ (177 mL)

Ages 12+ Years

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

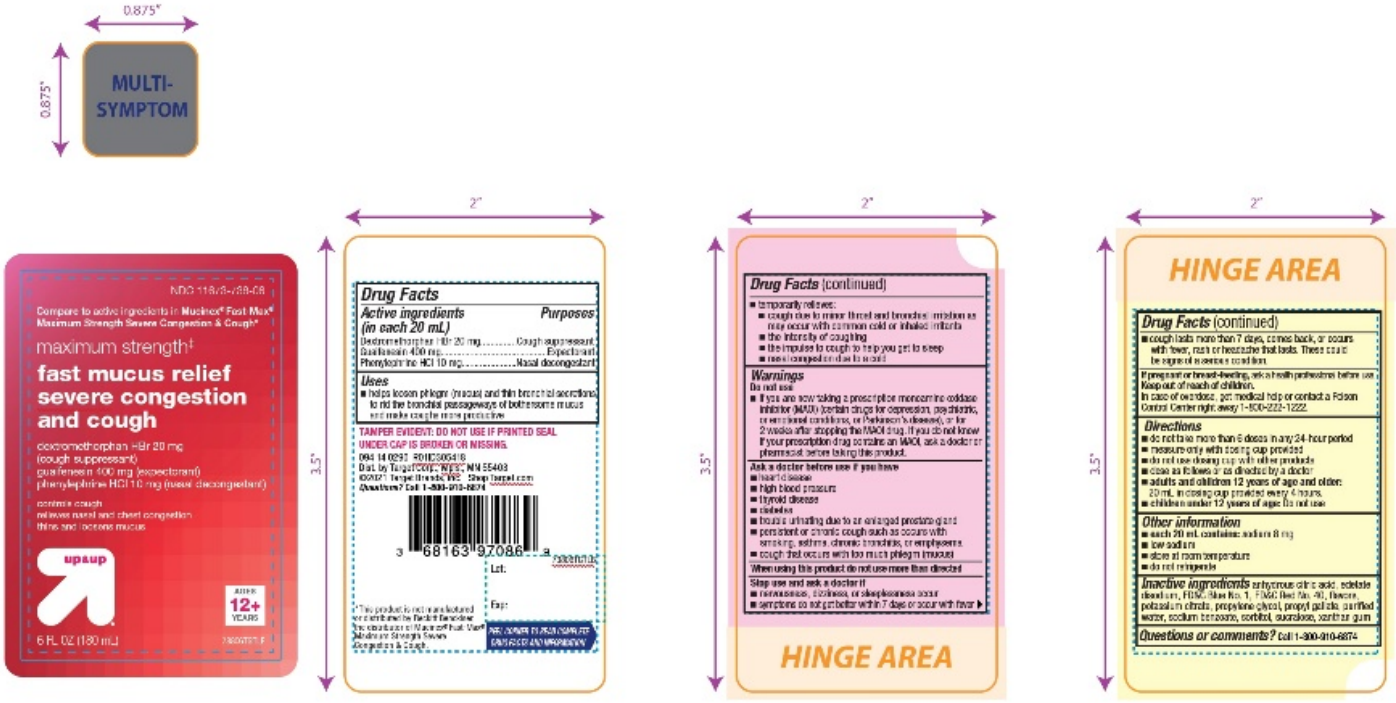
094 14 0290 R00 ID305418

Dist.by Target Corp., Mpls., MN 55403

Questions? Call 1-800-910-6874

‡Maximum Strength per 4 hour dose.

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



UP AND UP MAXIMUM STRENGTH FAST MUCUS RELIEF SEVERE CONGESTION AND COUGH

dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-738
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)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
propylene glycol (UNII: 6DC9Q167V3)	
propyl gallate (UNII: 8D4SNN7V92)	

water (UNII: 059QF0KO0R)	
sodium benzoate (UNII: OJ245FE5EU)	
sorbitol (UNII: 506T60A25R)	
sucralose (UNII: 96K6UQ3ZD4)	
xanthan gum (UNII: TTV12P4NEE)	

Product Characteristics

Color	BLUE (viscous liquid)	**Score**	
Shape		**Size**	
Flavor	FRUIT	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-738-06	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/10/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	11/10/2015	

Labeler - Target Corporation (006961700)

Revised: 11/2023 Target Corporation