

**UP AND UP COUGH PLUS CHEST CONGESTION- dextromethorphan hbr,
guaifenesin solution
Target Corporation**

Target Corporation Cough + Chest Congestion Drug Facts

Active ingredients (in each 20 mL)

Dextromethorphan HBr, USP 20 mg

Guaifenesin, USP 400 mg

Purposes

Cough suppressant

Expectorant

Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes

Warnings

Do not use

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

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

Stop use and ask a doctor if

cough lasts for more than 7 days, comes back, or is accompanied by fever, rash, or persistent headache. A persistent cough may be a sign 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
- keep dosing cup with product
- mL = milliliter
- this adult product is not intended for use in children under 12 years of age

age	dose
adults and children 12 years and over	20 mL every 4 hours
children under 12 years	do not use

Other information

- **each 20 mL contains:** sodium 13 mg
- store at 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

acetic acid, anhydrous citric acid, carboxymethylcellulose sodium, FD&C blue no. 1, FD&C red no. 40, flavor, glycerin, menthol, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol solution, sucralose, xanthan gum

Questions or comments?

1-888-547-7400

Package/Label Principal Display Panel

Compare to active ingredients in Robitussin® Maximum Strength Cough + Chest Congestion DM

Maximum Strength / Non-Drowsy

Cough & Chest Congestion

DM MAX

Dextromethorphan HBr (Cough Suppressant)

Guaifenesin (Expectorant)

- Relieves cough and mucus
- For adults
- Alcohol free

Ages 12+ Years

up&up™

Raspberry Menthol Flavor

8 FL OZ (236 mL)



Compare to active ingredients in Robitussin® Maximum Strength Cough + Chest Congestion DM*

Maximum Strength / Non-Drowsy

Cough & Chest Congestion DM MAX

Dextromethorphan HBr (Cough Suppressant)
Guaifenesin (Expectorant)

- Relieves cough and mucus
- For adults
- Alcohol free



8 FL OZ (236 mL)

Ages 12+ Years



Raspberry Menthol Flavor

Maximum Strength / Non-Drowsy

Cough & Chest Congestion DM MAX

Dextromethorphan HBr (Cough Suppressant)
Guaifenesin (Expectorant)



Maximum Strength / Non-Drowsy
Cough & Chest Congestion DM MAX
Dextromethorphan HBr (Cough Suppressant)
Guaifenesin (Expectorant)
up&up

Drug Facts

Active ingredients (in each 20 mL)
Dextromethorphan HBr, USP 20 mg.....Cough suppressant
Guaifenesin, USP 400 mg.....Expectorant

Purposes

- Uses**
- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
 - helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes

Warnings
Do not use 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**
- cough that occurs with too much phlegm (mucus)
 - cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis, or emphysema

Stop use and ask a doctor if cough lasts for more than 7 days, comes back, or is accompanied by fever, rash, or persistent headache. A persistent cough may be a sign 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
 - keep dosing cup with product
 - mL = milliliter
 - this adult product is not intended for use in children under 12 years of age

age	dose
adults and children 12 years and over	20 mL every 4 hours
children under 12 years	do not use

Drug Facts (continued)

Other information

- each 20 mL contains: sodium 13 mg
- store at 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients: acetic acid, anhydrous citric acid, carboxymethylcellulose sodium, FD&C blue no. 1, FD&C red no. 40, flavor, glycerin, menthol, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol solution, sucralose, xanthan gum

Questions or comments? 1-888-547-7400

*This product is not manufactured or distributed by Pfizer, distributor of Robitussin® Maximum Strength Cough + Chest Congestion DM.

DO NOT USE IF PRINTED NECKBAND IS BROKEN OR MISSING

GLUTEN FREE

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

Satisfaction guaranteed - Love it or your money back.

Distributed by Target Corporation Minneapolis, MN 55403 TM & © 2024 Target Brands, Inc.

NDC 11673-627-34



2B734 UW C2

UP AND UP COUGH PLUS CHEST CONGESTION

dextromethorphan hbr, guaifenesin solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-627
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

Inactive Ingredients

Ingredient Name	Strength
ACETIC ACID (UNII: Q40Q9N063P)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	RED	Score	
Shape		Size	
Flavor	FRUIT	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-627-34	1 in 1 CARTON	08/23/2018	
1		236 mL in 1 BOTTLE; Type 0: Not a Combination Product		

2	NDC:11673-627-26	1 in 1 CARTON	10/14/2022	04/30/2026
2		118 mL in 1 BOTTLE; 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	08/23/2018	

Labeler - Target Corporation (006961700)

Revised: 11/2024

Target Corporation