7 SELECT MAXIMUM STRENGTH- dextromethorphan hbr, guaifenesin solution
7-ELEVEN

7-ELEVEN Non-Drowsy Maximum Strength 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.

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Keep out of reach of children.

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

Directions

- shake well before using
- 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 7 mg
- store at room temperature. Do not refrigerate.
- Alcohol-free

Inactive ingredients

anhydrous citric acid, carboxymethylcellulose sodium, edetate disodium, FD&C Red No. 40, flavor, menthol, microcrystalline cellulose, povidone, propylene glycol, purified water, potassium citrate, sodium benzoate, sorbitol solution, sucralose, xanthan gum.

Questions or comments?

1-844-428-2538

Package/Label Principal Display Panel

Compare to Robitussin® Cough + Chest CONGESTION DM the active ingredients*

NDC# 10202-740-04

Non-Drowsy

Maximum Strength

Cough + Chest Congestion DM

Dextromethorphan HBr

Cough Suppressant

Guaifenesin

Expectorant

Relieves Chest Congestion Controls cough and mucus

QUALITY GUARANTEED

For Ages 12 & Over

4 FL OZ (118 mL)

DISTRIBUTED BY 7-ELEVEN, INC.

IRVING, TX 75063

WWW.7-ELEVEN.COM

SATISFACTION GUARANTEED 1-800-255-0711

IMPORTANT: Keep this carton for future reference on full labeling.

*This product is not manufactured or distributed by Pfizer Consumer Healthcare, owner of the registered trademark Robitussin ®Cough + Congestion DM.

7 SELECT MAXIMUM STRENGTH

dextromethorphan hbr, guaifenesin solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10202-740
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH)	DEXTROMETHORPHAN	20 mg

(DEXTROMETHORPHAN - UNII:7355X3ROTS)	HYDROBROMIDE	in 20 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL (UNII: 506T60A25R)	
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:10202-740-04	1 in 1 CARTON	03/25/2019	
1		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	03/25/2019	

Labeler - 7-ELEVEN (007347602)

Revised: 7/2024 7-ELEVEN