ADULT COUGH PLUS CHEST CONGESTION DM WITH HONEY-
dextromethorphan hbr, guaifenesin solution
WALGREEN COMPANY

Adult Cough + Chest Congestion DM with Honey

Drug Facts

Active ingredients (in each 20 ml)	***Purposes***
Dextromethorphan HBr, USP 20 mg	Cough suppressant
Guaifenesin, USP 400 mg	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 more than 7 days, comes back, or is accompanied by fever, 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 at 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 10 mg
- store at room temperature. Do not refrigerate.

Inactive ingredients

citric acid, carboxymethylcellulose sodium, edetate disodium, glycerin, honey, natural & artificial flavors, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate, sucralose, xanthan gum, zinc gluconate

Questions or comments?

1-866-467-2748

PRINCIPAL DISPLAY PANEL - 118 ml Bottle Label

Compare to active ingredients in Robitussin® Honey Cough + Chest Congestion DM MAX*

NDC 0363-7754-04

Adult

Cough + Chest

Congestion DM

With Honey

Dextromethorphan HBr (Cough Suppressant)
Guaifenesin (Expectorant)

Maximum Strength

DM MAX

- Controls Cough
- Relieves Chest Congestion
- Thins & Loosens Mucus
- Gluten Free
- No Artificial Colors
- No Added Alcohol

Taste the Real Honey

For Ages 12 & Over

4 FL OZ (118 ml)

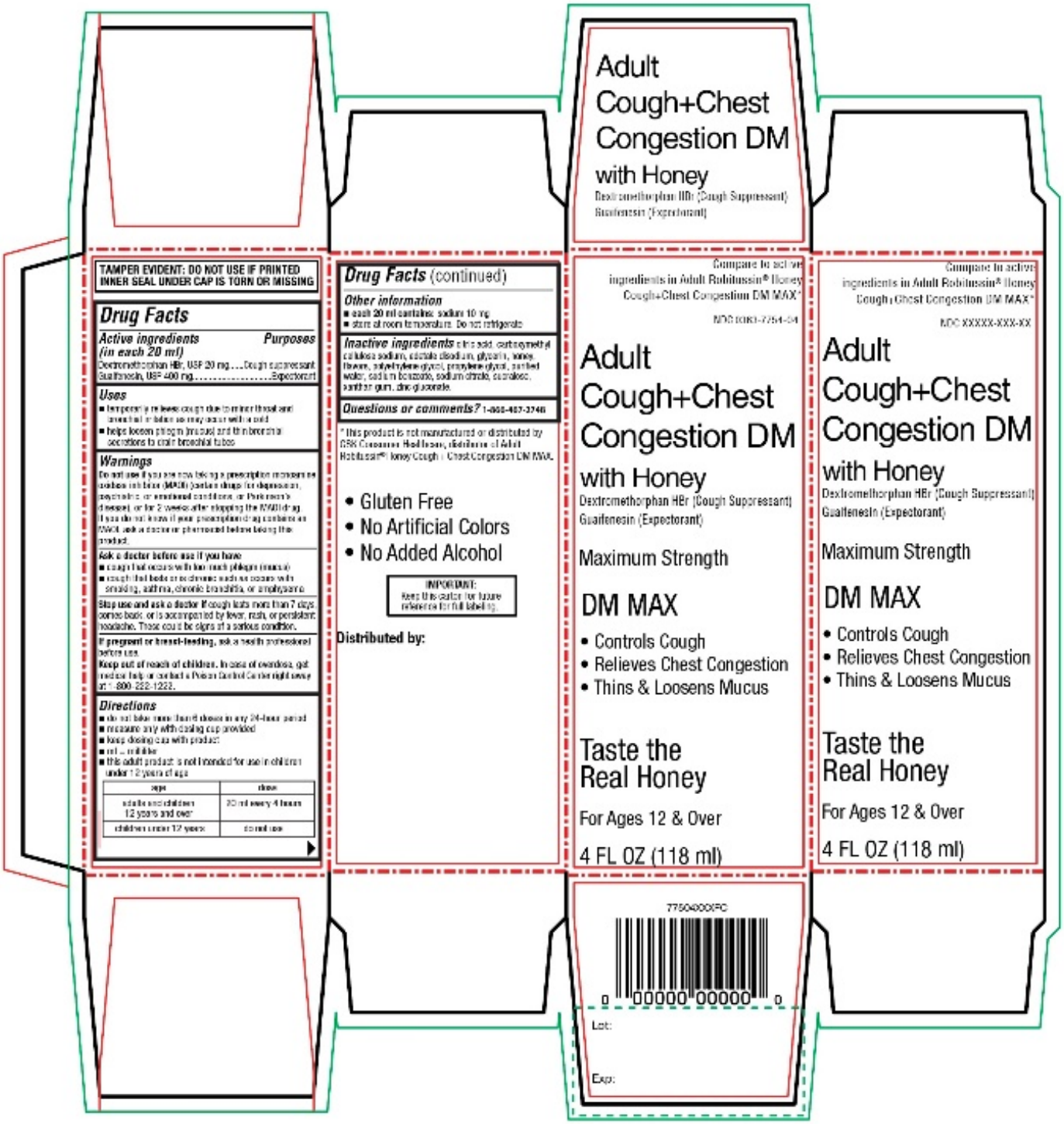
*This product is not manufactured or distributed by GSK Consumer Healthcare,

distributor of Adult Robitussin® Honey Cough + Chest Congestion DM MAX.

TAMPER EVIDENT: DON NOT USE IF PRINTED INNER SEAL UNDER CAP IS TORN OR MISSING

IMPORTANT: Keep this carton for future reference for full labelling.

Distributed by:



ADULT COUGH PLUS CHEST CONGESTION DM WITH HONEY

dextromethorphan hbr, guaifenesin solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-7754
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
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
HONEY (UNII: Y9H1V576FH)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ZINC GLUCONATE (UNII: U6WSN5SQ1Z)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-7754-04	1 in 1 CARTON	02/05/2024	
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	02/05/2024	

Labeler - WALGREEN COMPANY (008965063)