

**TUSSIN DM COUGH AND CHEST CONGESTION ADULT- dextromethorphan hbr,  
guaifenesin liquid  
P & L Development, LLC**

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**Drug Facts**

**Active ingredients (in each 10 mL)**

Dextromethorphan HBr 20 mg

Guaifenesin 200 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 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 (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
- this adult product is not intended for use in children under 12 years of age
- adults and children 12 years and over: 10 mL every 4 hours
- children under 12 years: do not use

### **Other information**

- store between 20-25°C (68-77°F). Do not refrigerate.

### **Inactive ingredients**

citric acid, FD&C red# 40, flavor, glucose, glycerin, high fructose corn syrup, menthol, purified water, saccharin sodium, sodium benzoate

### **Questions or comments?**

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

### **Principal Display Panel**

Compare to the active ingredients in Robitussin Cough + Chest Congestion DM\*

adult tussin cough & chest congestion DM

Dextromethorphan HBr 20 mg

Guaifenesin 200 mg

cough suppressant

expectorant

relieves:

- Cough
- chest congestion
- mucus

for ages 12+

alcohol free

fl oz (mL)

\*This product is not manufactured or distributed by Haleon group of companies, distributor of Robitussin Cough+Chest Congestion DM.

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

Manufactured by:

**PL Developments**

11865 S. Alameda St

Lynwood, CA 90262

**Package Label**

Compare to the active ingredients in Robitussin Cough+Chest Congestion DM\* NDC 49580-0385-2

**ready in case**

**adult tussin cough & chest congestion DM**

**Dextromethorphan HBr 20 mg**  
**Guaifenesin 200 mg**  
cough suppressant  
expectorant

relieves:  
• cough  
• chest congestion  
• mucus

for ages 12+  
alcohol free

4 fl oz (118 mL)

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**Active ingredients (in each 10 mL)**  
Dextromethorphan HBr 20 mg.....Cough suppressant  
Guaifenesin 200 mg.....Expectorant

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

Manufactured by:  
**PL Developments**  
11865 S. Alameda St  
Lynwood, CA 90262

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**Drug Facts** (continued under label) **PEEL HERE** →

**Drug Facts** (continued)

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**READYinCASE Adult Tussin Cough & Chest Congestion DM**

**TUSSIN DM COUGH AND CHEST CONGESTION ADULT**

dextromethorphan hbr, guaifenesin liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:49580-0385
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 10 mL
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 10 mL

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>DEXTROSE</b> (UNII: IY9XDZ35W2)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>HIGH FRUCTOSE CORN SYRUP</b> (UNII: XY6UN3QB6S)	
<b>MENTHOL</b> (UNII: L7T10EIP3A)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SACCHARIN SODIUM</b> (UNII: SB8ZUX40TY)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:49580-0385-4	1 in 1 BOX	03/31/2015	
1		118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:49580-0385-2	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/31/2015	
3	NDC:49580-0385-8	1 in 1 BOX	03/31/2015	
3		237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M012	03/31/2015	

**Labeler** - P & L Development, LLC (101896231)