

**TUSSIN DM MAX- dextromethorphan hbr, guaifenesin solution**  
**Wal-Mart Stores Inc**

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**Equate 44-030**

***Active ingredients (in each 20 mL)***

Dextromethorphan HBr 20 mg  
Guaifenesin 400 mg

***Purpose***

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)
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema

**Stop use and ask a doctor if**

cough persists more than 7 days, tends to recur, or is accompanied by a 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.

***Directions***

- **do not take more than directed**
- do not take more than 6 doses in any 24-hour period
- mL = milliliter
- only use the dose cup provided
- adults and children 12 years and over: 20 mL in dosing cup provided every 4 hours
- children under 12 years: do not use

***Other information***

- **each 20 mL contains:** sodium 16 mg
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

***Inactive ingredients***

anhydrous citric acid, FD&C blue #1, FD&C red #40, flavors, glycerin, high fructose corn syrup, microcrystalline cellulose, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium chloride, sodium citrate dihydrate, sorbitol, sucralose, xanthan gum

***Questions or comments?***

**1-888-287-1915**

**Principal Display Panel**

NDC 49035-930-19

**equate™**

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

**Tussin DM  
Max**

**Cough & Chest  
Congestion DM**

**Oral Solution**

**Dextromethorphan HBr -  
Cough Suppressant  
Guaifenesin - Expectorant**

**Maximum Strength**

- Controls cough
- Relieves chest congestion
- Thins & loosens mucus

**Ages 12+**

Raspberry Flavor  
Dosage cup included

**8 FL OZ (237 mL)**

**TAMPER EVIDENT: DO NOT USE IF  
PRINTED NECK WRAP IS BROKEN  
OR MISSING**

**PARENTS:**

Learn about teen medicine abuse  
[www.StopMedicineAbuse.org](http://www.StopMedicineAbuse.org)

Satisfaction guaranteed –  
Or we'll replace it or give you  
your money back. For questions  
or comments or to report an  
undesired reaction or side effect,  
please call **1-888-287-1915**.

**DISTRIBUTED BY: Walmart Inc.,  
Bentonville, AR 72716**

**\*\*This product is not manufactured or distributed  
by Haleon US Holdings LLC, owner of the registered  
trademark Robitussin® Maximum Strength  
Cough+Chest Congestion DM.**

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Equate 44-030

## TUSSIN DM MAX

dextromethorphan hbr, guaifenesin solution

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-930
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)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	red (Maroon)	Score	
Shape		Size	
Flavor	BERRY, MENTHOL	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-930-19	1 in 1 CARTON	08/15/2018	
1	NDC:49035-930-00	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
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Category	Citation	Date	Date
OTC Monograph Drug	M012	08/15/2018	

**Labeler** - Wal-Mart Stores Inc (051957769)

**Establishment**

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	manufacture(49035-930) , pack(49035-930)

Revised: 1/2025 Wal-Mart Stores Inc