SDA LABORATORIES TUSSIN DM- dextromethorphan hbr, guaifenesin liquid
SDA Laboratories, Inc.

SDA-TussinDM 565

per 5 mL teaspoonful

Dextromethorphan HBr - 10 mg

Guaifenesin - 100 mg

Purpose

Cough Suppressant

Expectorant

Uses

- temporarilyrelievescoughduetominorthroatand bronchial irritation
- helps loosen phlegm (mucus)
- helps thin bronchial secretions to make coughs more productive

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.

Do not useif you have ever had an allergic reaction to any of the ingredients in this product.

Ask a doctor before use if you have

- a cough with too much phlegm (mucus)
- a 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 headache that lasts. 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

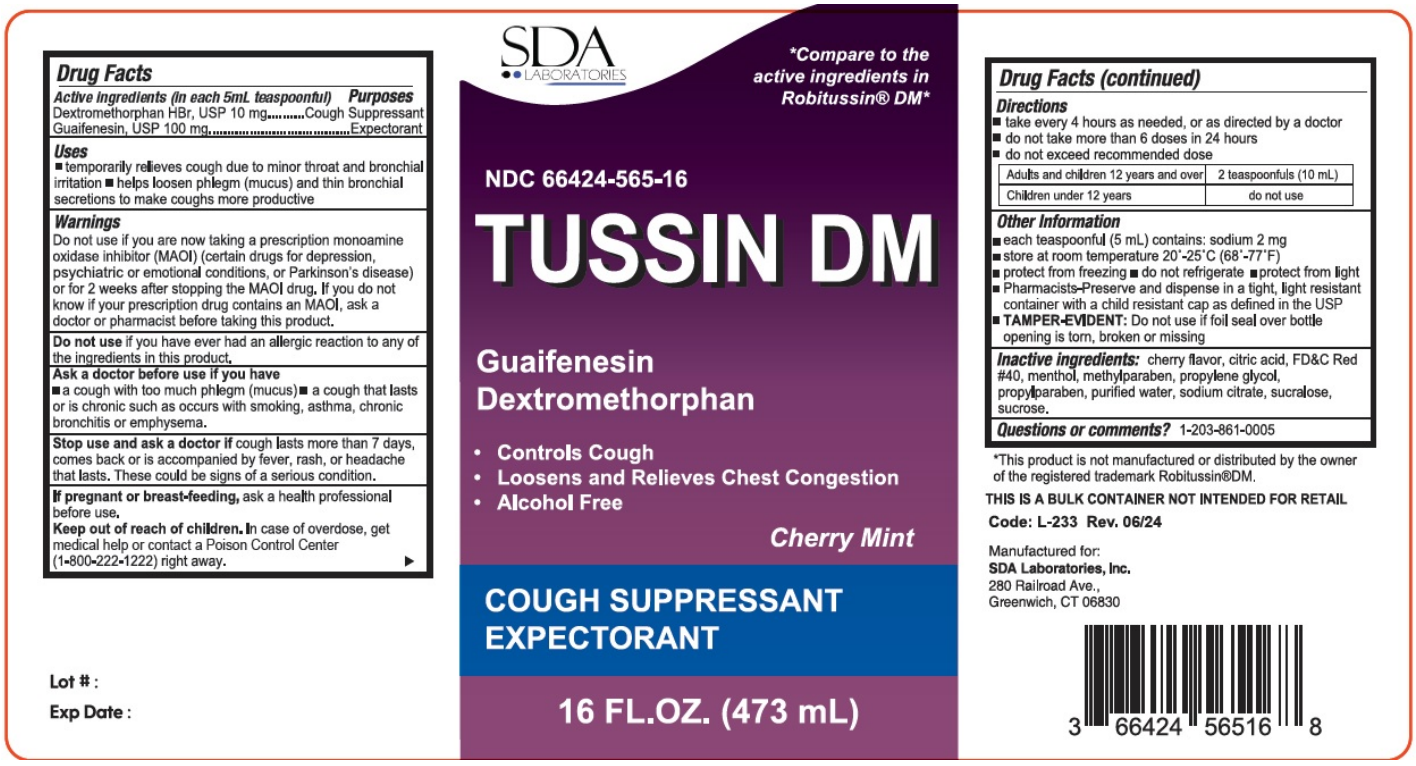
- take every 4 hours as needed, or as directed by a doctor
- do not take more than 6 doses in 24 hours
- do not exceed recommended dose

Adults and children 12 years and over	2 teaspoonfuls (10 mL)

Children under 12 years	do not use

Inactive ingredients: artificial and natural cherry flavor, citric acid, FD&C Red #40, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate, sucralose, sucrose

Questions or Comments? 1-203-861-0005



SDA LABORATORIES TUSSIN DM

dextromethorphan hbr, guaifenesin liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66424-565
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 5 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 5 mL

Inactive Ingredients	
Ingredient Name	**Strength**
MENTHOL (UNII: L7T10EIP3A)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

FD&C RED NO. 40 (UNII: WZB9127XOA)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SUCROSE (UNII: C151H8M554)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66424-565-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/01/2024	

Labeler - SDA Laboratories, Inc. (948067889)

Revised: 12/2024 SDA Laboratories, Inc.