

TUSSIN DM- dextromethorphan hbr guaifenesin solution
Strategic Sourcing Services LLC

Foster and Thrive 44-073

Active ingredients (in each 20 mL)

Dextromethorphan HBr 20 mg
Guaifenesin 200 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 every 4 hours
- children under 12 years: do not use

Other information

- **each 20 mL contains:** sodium 8 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, flavors, glycerin, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate dihydrate, sucralose, xanthan gum

Questions or comments?

Call **833-358-6431** Monday to Friday 9:00am to 7:00pm EST

Principal display panel

NDC 70677-1261-1

*COMPARE TO
THE ACTIVE INGREDIENTS IN
ROBITUSSIN® SUGAR-FREE DYE-FREE
COUGH+CHEST CONGESTION DM

Foster & Thrive™

Tussin DM

Dextromethorphan HBr
Guaifenesin
COUGH SUPPRESSANT
EXPECTORANT

Ages 12 Years and Over**Oral Solution**

Sugar Free

Dye Free

Cool Mint Flavored

- **Controls Cough**
- **Relieves Chest Congestion**
- **Thins & Loosens Mucus**

DOSING CUP

INCLUDED

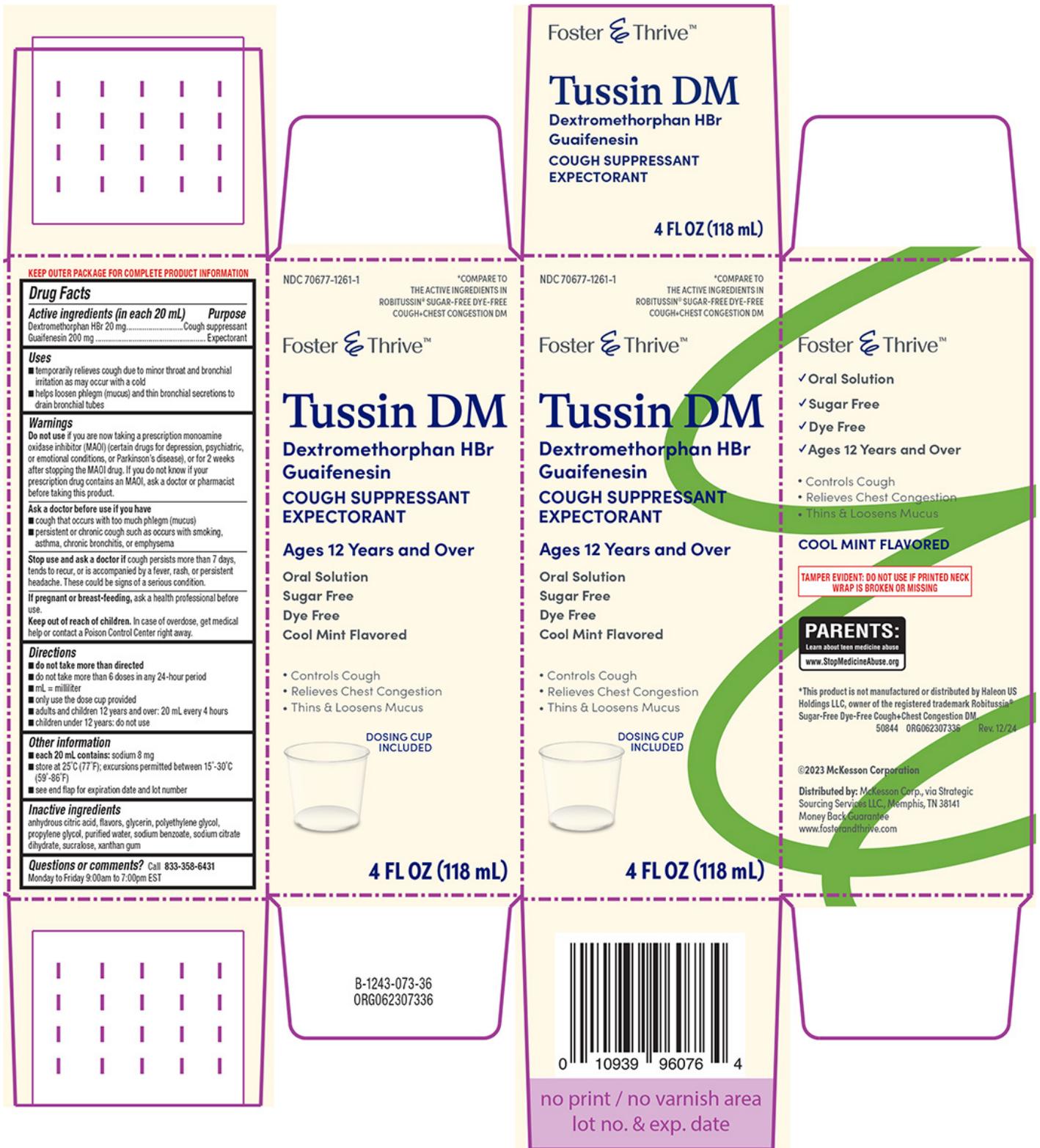
4 FL OZ (118 mL)

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

***This product is not manufactured or distributed by Haleon US
Holdings LLC, owner of the registered trademark Robitussin®
Sugar-Free Dye-Free Cough+Chest Congestion DM.
50844 ORG062307336 Rev. 12/24**

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Sourcing Services LLC., Memphis, TN 38141
Money Back Guarantee
www.fosterandthrive.com**



Foster and Thrive 44-073

TUSSIN DM

dextromethorphan hbr guaifenesin solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70677-1261
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	200 mg in 20 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0K00R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	MENTHOL, MINT	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70677-1261-1	1 in 1 CARTON	02/26/2025	
1		118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:70677-1261-2	1 in 1 CARTON	02/26/2025	
2		237 mL in 1 BOTTLE, PLASTIC; 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	02/26/2025	

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	manufacture(70677-1261) , pack(70677-1261)

Revised: 1/2026

Strategic Sourcing Services LLC