GOODSENSE TUSSIN CF MULTI-SYMPTOM COLD, RASPBERRY-
dextromethorphan hbr, guaifenesin, phenylephrine hcl liquid
Geiss, Destin & Dunn Inc.

GoodSense Tussin CF Multi-Symptom Cold, Raspberry

Active ingredients (in each 5 mL or 1 teaspoon) Purposes
Dextromethorphan HBr, 10 mg..... Cough Suppressant
Guaifenesin, 100 mg..... Expectorant
Phenylephrine HCl, 5 mg..... Nasal Decongestant

Purposes
Cough Suppressant
Expectorant
Nasal Decongestant

Uses

• helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes temporarily relieves these symptoms occurring with a cold:

• nasal congestion • cough due to minor throat and bronchial irritation

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 • heart disease • high blood pressure
• thyroid disease • diabete s • trouble urinating due to an enlarged prostate gland
• cough that occurs with too much phlegm (mucus) • cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema

Ask a doctor or pharmacist before use if you are taking any other oral nasal decongestant or stimulant.

Do not exceed recommended dosage.

Stop use and ask a doctor if • you get nervous, dizzy or sleepless
• symptoms do not get better within 7 days or are accompanied by fever • cough lasts more than 7 days, comes back, or is accompanied by fever, rash, or persistant 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 6 doses in any 24 hour period • this adult strength product is not intended for use in children under 12 years of age • measure

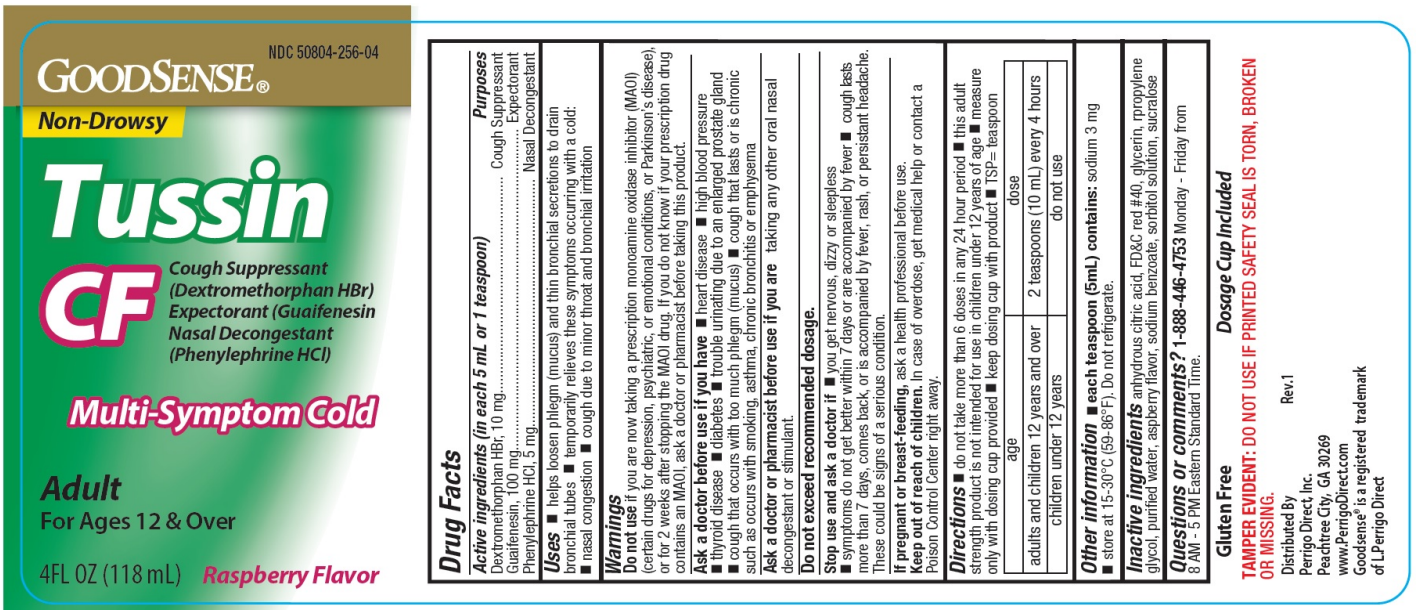
only with dosing cup provided • keep dosing cup with product • TSP= teaspoon

age	dose
adults and children 12 years and over	2 teaspoons (10 mL) every 4 hours
children under 12 years	do not use

Other information
- **each teaspoon (5mL) contains:** sodium 3 mg
- store at 15-30°C (59-86°F). Do not refrigerate.

Inactive ingredients anhydrous citric acid, FD&C red #40, glycerin, propylene glycol, purified water, raspberry flavor, sodium benzoate, sorbitol solution, sucralose

Questions or comments? 1-888-446-4753 Monday - Friday from 8 AM - 5 PM Eastern Standard Time.



GOODSENSE TUSSIN CF MULTI-SYMPTOM COLD, RASPBERRY

dextromethorphan hbr, guaifenesin, phenylephrine hcl liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50804-256
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 5 mL
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
WATER (UNII: 059QF0KO0R)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
SORBITOL SOLUTION 70% (UNII: 8KW3E207O2)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	RASPBERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50804-256-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/05/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	11/05/2024	

Labeler - Geiss, Destin & Dunn Inc. (076059836)

Registrant - Geiss, Destin & Dunn Inc. (076059836)

Revised: 9/2025 Geiss, Destin & Dunn Inc.