TUKOL COUGH AND CONGESTION- dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride syrup
Genomma Lab USA

Tukol ®Cough & Congestion Syrup

Drug Facts

Active Ingredient (in each 5 mL or 1 teaspoon)	**Purpose**
Dextromethorphan HBr, 10 mg	Cough Suppressant
Guaifenesin, 100 mg	Expectorant
Phenylephrine HCl, 5 mg	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 useif 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
- diabetes
- 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 aretaking 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 (5 mL) contains:** sodium 3 mg
- store at 15-30°C (59-86°F). Do not refrigerate.

Inactive ingredients

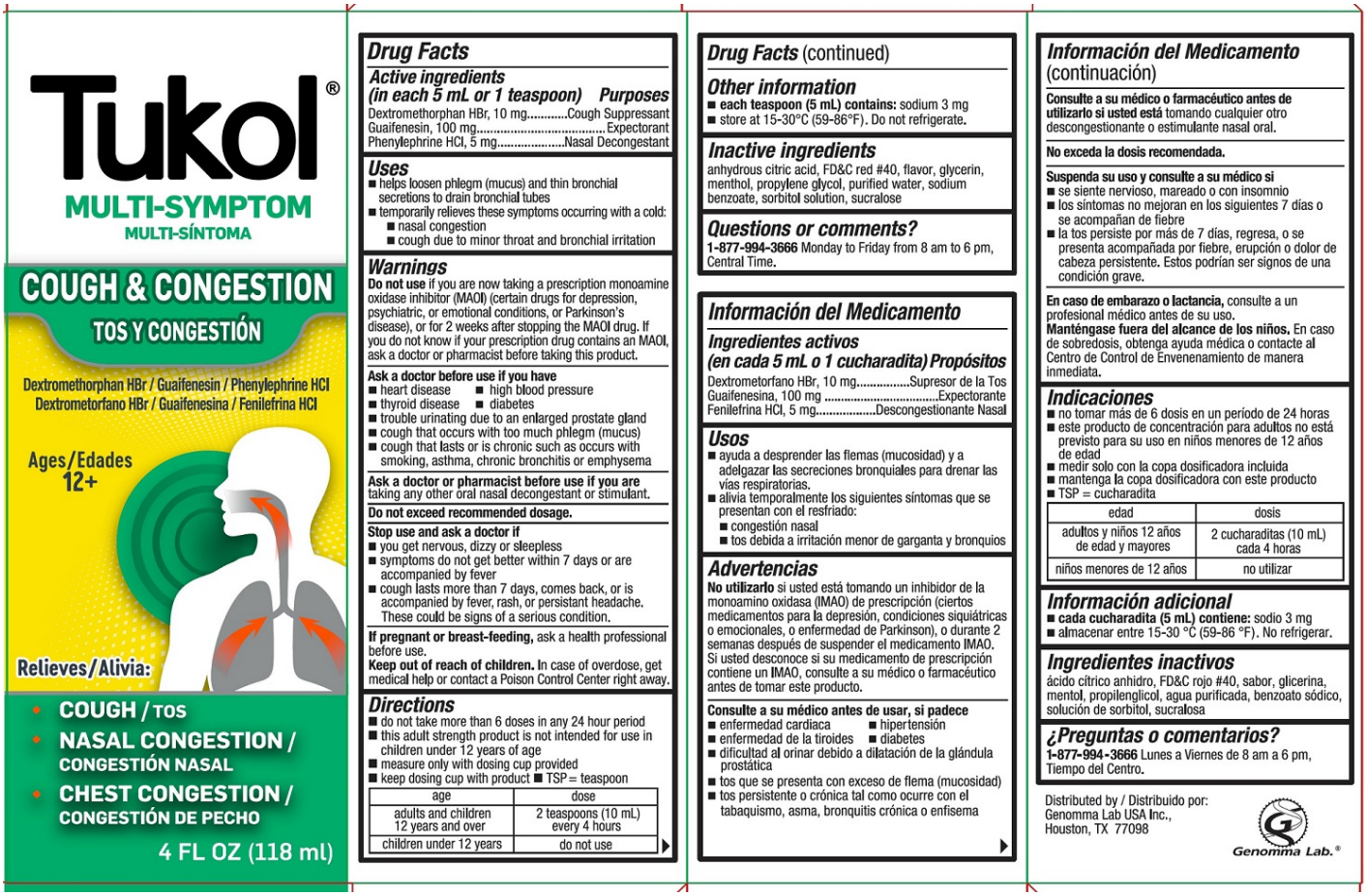
anhydrous citric acid, FD&C red #40, flavor, glycerin, menthol, propylene glycol, purified water, sodium benzoate, sorbitol solution, sucralose

Questions or comments?

1-877-994-3666 Monday to Friday from 8 am to 6 pm, Central Time.

Distributed by:
Genomma Lab USA Inc.,
Houston, TX 77027

PRINCIPAL DISPLAY PANEL - 118 ml Bottle Carton



Tukol ®

MULTI-SYMPTOM

COUGH & CONGESTION

Dextromethorphan HBr / Guaifenesin / Phenylephrine HCl

Ages
12+

Relieves:

- COUGH
- NASAL CONGESTION
- CHEST CONGESTION

4 FL OZ (118 ml)

TUKOL COUGH AND CONGESTION

dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride syrup

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50066-510
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
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHOL (UNII: L7T10EIP3A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50066-510-04	1 in 1 CARTON	06/07/2020	
1		118 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	06/07/2020	

Labeler - Genomma Lab USA (832323534)

Registrant - Genomma Lab USA (832323534)