VICKS CHILDRENS COUGH CONGESTION- phenylephrine hcl, dextromethorphan hbr, guaifenesin liquid
The Procter & Gamble Manufacturing Company

VICKS ®

children's Cough Congestion

Drug Facts

Active ingredients (in each 15 mL)

Dextromethorphan HBr 10 mg

Guaifenesin 200 mg

Phenylephrine HCl 5 mg

Purpose

Cough suppressant

Expectorant

Nasal decongestant

Uses

temporarily relieves common cold symptoms:

- nasal congestion
- cough due to minor throat and bronchial irritation
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and 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.

Ask a doctor before use if you have

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to enlarged prostate gland
- cough that occurs with too much phlegm (mucus)

- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis or emphysema
- a sodium-restricted diet

When using this product, do not use more than directed.

Stop use and ask a doctor if
- you get nervous, dizzy or sleepless
- symptoms do not improve within 7 days or occur with a fever
- cough persists for more than 7 days, comes back or occurs with 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
- take only as directed
- only use the dose cup provided
- do not exceed 4 doses per 24 hrs

adults & children 12 yrs & over	30 mL every 4 hrs
children 6 to under 12 yrs	15 mL every 4 hrs
children 4 to under 6 yrs	7.5 mL every 4 hrs
children under 4 yrs	do not use

Other information
- **each 15 mL contains:**sodium 44 mg
- store at no greater than 25ºC.

Inactive ingredients

anhydrous citric acid, flavor, glycerin, propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium chloride, sodium citrate, sorbitol solution, sucralose

Questions?

1-800-362-1683

TAMPER EVIDENT: Do not use if printed safety seal under cap is missing or damaged.

DIST. BY: PROCTER & GAMBLE, CINCINNATI, OH 45202

Made in Canada

PRINCIPAL DISPLAY PANEL - 177 ml Bottle Label

VICKS ®

children's

Cough

Congestion

Non-Drowsy

Phenylephrine HCl - nasal decongestant
Dextromethorphan HBr - cough suppressant
Guaifenesin - expectorant

FREE of:

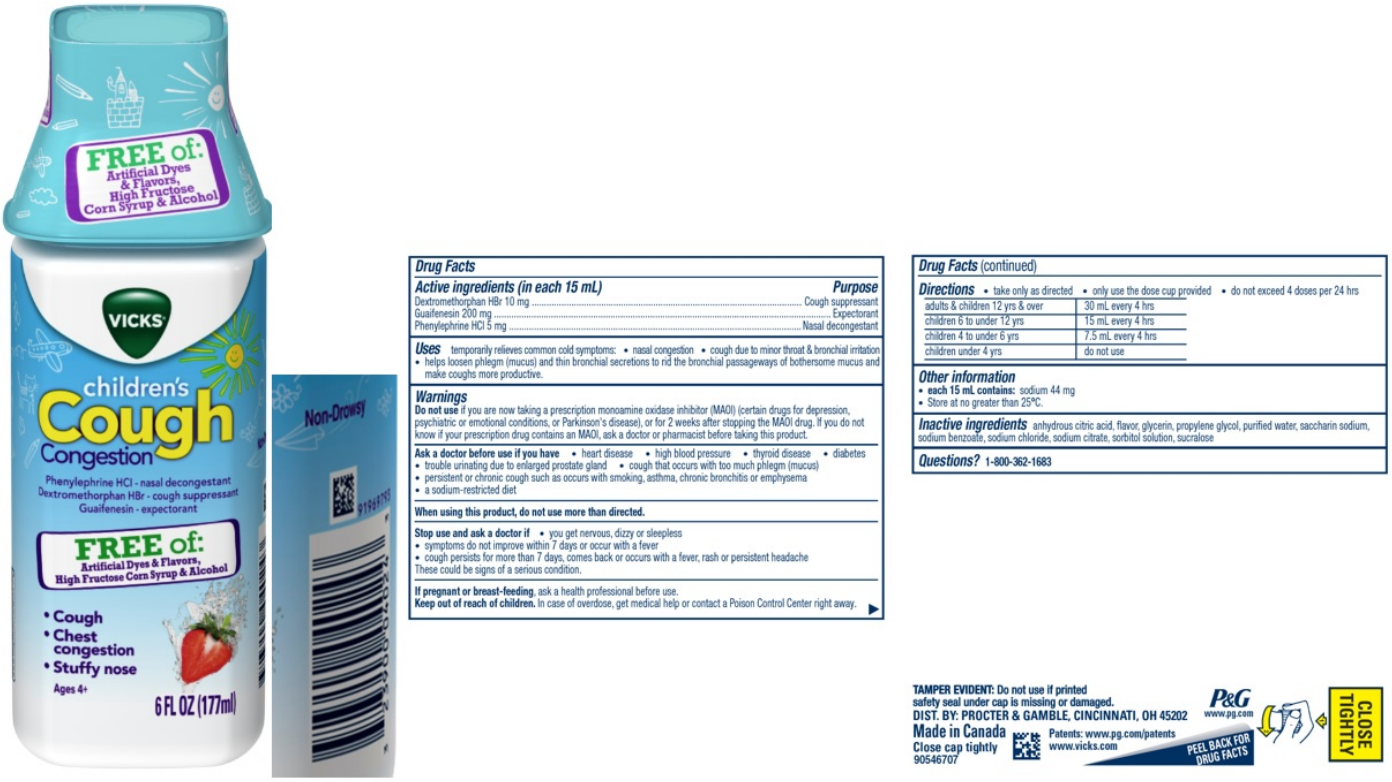
Artificial Dyes & Flavors,

High Fructose Corn Syrup & Alcohol

- Cough
- Chest congestion
- Stuffy nose

Ages 4+

6 FL OZ (177 ml)



VICKS CHILDRENS COUGH CONGESTION

phenylephrine hcl, dextromethorphan hbr, guaifenesin liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69423-982
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 15 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 15 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

SUCRALOSE (UNII: 96K6UQ3ZD4)	
SORBITOL (UNII: 506T60A25R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Product Characteristics			
Color	white (Clear)	**Score**	
Shape		**Size**	
Flavor	STRAWBERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-982-06	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/22/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	04/22/2019	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Establishment

Name	Address	ID/FEI	Business Operations
SPERA NEXUS INC		097722284	api manufacture(69423-982)

Revised: 10/2023 The Procter & Gamble Manufacturing Company