VICKS CHILDRENS COUGH CONGESTION NIGHT- phenylephrine hydrochloride and diphenhydramine hydrochloride liquid
The Procter & Gamble Manufacturing Company

VICKS ®
children's Cough Congestion Night

Drug Facts

Active ingredients (in each 15 mL)

Diphenhydramine HCl 12.5 mg

Phenylephrine HCl 5 mg

Purpose

Antihistamine/Cough suppressant
Nasal decongestant

Uses

temporarily relieves common cold symptoms:
- nasal congestion
- runny nose & sneezing
- cough due to minor throat and bronchial irritation

Warnings

Do not use
- with any other product containing diphenhydramine, even one used on skin
- to make a child sleepy
- 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
- glaucoma
- trouble urinating due to enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma, or emphysema

- a sodium-restricted diet

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers

When using this product

- **do not use more than directed.**
- excitability may occur, especially in children
- marked drowsiness may occur
- alcohol, sedatives, and tranquilizers may increase drowsiness
- avoid alcoholic drinks
- be careful when driving a motor vehicle or operating machinery

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.

Other information

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

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	do not use unless directed by a doctor
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, polysorbate 20, 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 **Night**

Phenylephrine HCl - nasal decongestant
Diphenhydramine HCl - antihistamine/cough suppressant

FREEof:

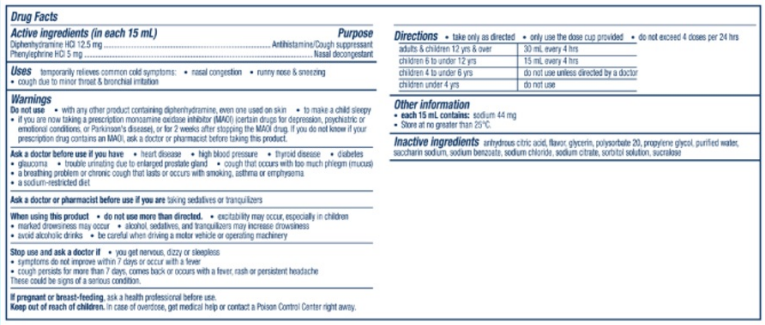
Artificial Dyes & Flavors,

High Fructose Corn Syrup & Alcohol

- Cough
- Runny nose
- Stuffy nose
- Sneezing

Ages 6+

6 FL OZ (177 ml)



VICKS CHILDRENS COUGH CONGESTION NIGHT

phenylephrine hydrochloride and diphenhydramine hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37000-712
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	12.5 mg in 15 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL (UNII: 506T60A25R)	

Product Characteristics

Color	white (Clear)	**Score**	
Shape		**Size**	
Flavor	GRAPE	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37000-712-06	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)

Revised: 10/2024 The Procter & Gamble Manufacturing Company