

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 25°C.

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 25°C.

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)



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 that irritates throat & 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 the rash-prone illness (measles) • a breathing problem or chronic cough that lasts or 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 • medical conditions 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.	

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
 Patents: www.pg.com/patents
 Close cap tightly
 91969776

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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: 10/2023

The Procter & Gamble Manufacturing Company