

**VICKS NYQUIL CHILDRENS COLD AND COUGH- chlorpheniramine maleate,
dextromethorphan hbr liquid
The Procter & Gamble Manufacturing Company**

VICKS®
NyQuil™
children's COLD & COUGH

Drug Facts

Active ingredients (in each 15 mL)

Chlorpheniramine maleate 2 mg
Dextromethorphan HBr 15 mg

Purpose

Antihistamine
Cough suppressant

Uses

temporarily relieves cold symptoms:

- cough due to minor throat & bronchial irritation
- sneezing
- runny nose

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.
- to make a child sleepy

Ask a doctor before use if you have

- glaucoma
- cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma, chronic bronchitis or emphysema
- trouble urinating due to enlarged prostate gland

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

- excitability may occur, especially in children
- marked drowsiness may occur
- avoid alcoholic drinks
- be careful when driving a motor vehicle or operating machinery
- alcohol, sedatives, & tranquilizers may increase drowsiness

Stop use and ask a doctor if

- cough lasts more than 1 week, tends to recur, or is accompanied by 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 6 hrs
children 6 to under 12 yrs	15 mL every 6 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 13 mg
- Store at no greater than 25°C

Inactive ingredients

citric acid, FD&C Red No. 40, flavor, glycerin, propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium citrate, sorbitol, sucralose, xanthan gum

Questions?

1-800-362-1683

TAMPER EVIDENT: Do not use if printed shrinkband is broken or missing.

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

PRINCIPAL DISPLAY PANEL - 236 ml Bottle Label

VICKS®

NyQuil™
children's

COLD & COUGH

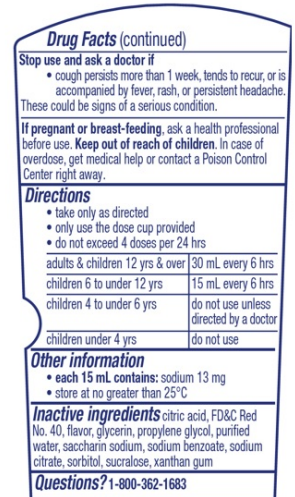
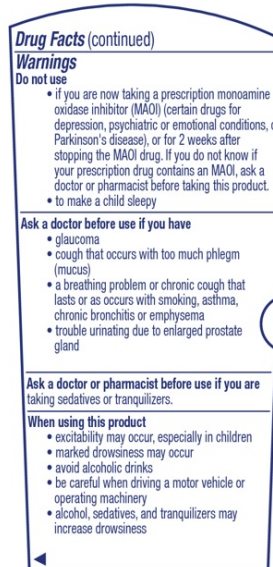
Chlorpheniramine maleate, Dextromethorphan HBr

Sneezing, Runny Nose
Cough

Free of Alcohol & Acetaminophen

8 FL OZ (236 mL)

AGES 6+



VICKS NYQUIL CHILDRENS COLD AND COUGH

chlorpheniramine maleate, dextromethorphan hbr liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69423-898
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	2 mg in 15 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	15 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
GLYCERIN (UNII: PDC6A3C0OX)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SORBITOL (UNII: 506T60A25R)	

Product Characteristics

Color	red	Score	
Shape		Size	
Flavor	BERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-898-08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/07/2020	06/06/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	05/07/2020	06/06/2025

Revised: 10/2023

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