

VICKS NYQUIL COLD AND FLU- acetaminophen, dextromethorphan hydrobromide, and doxylamine succinate liquid
The Procter & Gamble Manufacturing Company

VICKS® NyQuil COLD & FLU

Drug Facts

Active ingredients (in each 30 mL)

Acetaminophen 650 mg

Dextromethorphan HBr 30 mg

Doxylamine succinate 12.5 mg

Purpose

Pain reliever/fever reducer

Cough suppressant

Antihistamine

Uses

temporarily relieves common cold/flu symptoms:

- cough due to minor throat and bronchial irritation
- sore throat
- headache
- minor aches and pains
- fever
- runny nose and sneezing

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take

- more than 4 doses in 24 hours, which is the maximum daily amount for this product
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Allergy Alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening • blisters • rash

If a skin reaction occurs, stop use and seek medical help right away

Sore throat warning: If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- 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

- liver disease
- 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
- taking the blood thinning drug warfarin

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, and tranquilizers may increase drowsiness

Stop use and ask a doctor if

- pain or cough gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- cough comes back or occurs with rash or headache that lasts.

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. Quick medical attention is critical for adults as well as for children even if you do not

notice any signs or symptoms.

Directions

- take only as directed
- only use the dose cup provided
- do not exceed 4 doses per 24 hrs

adults & children 12 yrs & over
children 4 to under 12 yrs
children under 4 yrs

30 mL every 6 hrs
ask a doctor
do not use

Other information

- **each 30 mL contains:**sodium 24 mg
- store at no greater than 25°C and do not refrigerate

Inactive ingredients

alcohol, citric acid, FD&C Blue No. 1, 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™

COLD & FLU

Acetaminophen, Doxylamine Succinate, Dextromethorphan HBr

- **Aches, Fever & Sore Throat**
- **Sneezing, Runny Nose**
- **Cough**

Nighttime Relief

Alcohol 10%

8 FL OZ (236 ml)



Drug Facts (continued)

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 4 to under 12 yrs ask a doctor
 children under 4 yrs do not use

Other information

- each 30 mL contains: sodium 24 mg
- store at no greater than 25°C and do not refrigerate

Inactive ingredients alcohol, citric acid, FD&C Blue No. 1, FD&C Red No. 40, flavor, glycerin, propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium citrate, sorbitol, sucralose, xanthan gum

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Drug Facts (continued)

bronchitis or emphysema
 • trouble urinating due to enlarged prostate gland

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VICKS NYQUIL COLD AND FLU

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69423-791
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg in 30 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg in 30 mL
DOXYLAMINE SUCCINATE (UNII: V9BI9B5Y12) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	12.5 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SORBITOL (UNII: 506T60A25R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALCOHOL (UNII: 3K9958V90M)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

XANTHAN GUM (UNII: TTV12P4NEE)

Product Characteristics

Color	red	Score	
Shape		Size	
Flavor	CHERRY, MENTHOL	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-791-08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2022	
2	NDC:69423-791-12	354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2022	
3	NDC:69423-791-24	2 in 1 PACKAGE, COMBINATION	01/01/2022	
3	NDC:69423-791-12	354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:69423-791-36	3 in 1 PACKAGE, COMBINATION	01/01/2022	
4	NDC:69423-791-12	354 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	01/01/2022	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 12/2025

The Procter & Gamble Manufacturing Company