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

VICKS ® NyQuil™ Severe COLD & FLU Liquicaps

Drug Facts

Active ingredients (in each LiquiCap)

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Doxylamine succinate 6.25 mg

Phenylephrine HCl 5 mg

Purpose

Pain reliever/fever reducer

Cough suppressant

Antihistamine

Nasal decongestant

Uses

temporarily relieves common cold/flu symptoms:

- nasal congestion
- sinus congestion & pressure
- cough due to minor throat & bronchial irritation
- cough to help you sleep
- minor aches & pains
- headache
- fever
- sore throat
- runny nose & sneezing
- reduces swelling of nasal passages
- temporarily restores freer breathing through the nose
- promotes nasal and/or sinus drainage

Warnings

Liver warning:

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

• more than 8 LiquiCaps 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
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- 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

- **do not use more than directed**
- 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

- you get nervous, dizzy or sleepless
- pain, nasal congestion, 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 • do not exceed 8 LiquiCaps per 24 hrs

adults & children 12 yrs & over	2 LiquiCaps with water every 4 hrs
children 4 to under 12 yrs	ask a doctor
children under 4 yrs	do not use

Other information

- do not exceed 25ºC

Inactive ingredients

FD&C Blue No. 1, gelatin, glycerin, pharmaceutical ink*, polyethylene glycol, povidone, propylene glycol, purified water, sorbitol sorbitan solution *May contain this ingredient

Questions?

1-800-362-1683

TAMPER EVIDENT: This package is safety-sealed & child resistant. Use only if blisters are intact. If difficult to open, use scissors.

Made in Canada

DIST. BY
PROCTER & GAMBLE,
CINCINNATI OH 45202

PRINCIPAL DISPLAY PANEL - 24 LiquiCap Carton

MAX

STRENGTH

VICKS®

NyQuil™

SEVERE

COLD & FLU

Acetaminophen, Phenylephrine HCl, Doxylamine Succinate **,**Dextromethorphan HBr

Headache, Fever, Sore Throat, Minor Aches & Pains
Nasal Congestion, Sinus Pressure
Sneezing, Runny Nose
Cough

Nighttime Relief

24 LIQUICAPS™

VICKS NYQUIL SEVERE COLD AND FLU

acetaminophen, phenylephrine hydrochloride, dextromethorphan hydrobromide, and doxylamine succinate capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37000-518
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
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	

Product Characteristics

Color	green	**Score**	no score
Shape	BULLET	**Size**	16mm
Flavor		**Imprint Code**	NS
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37000-518-24	2 in 1 CARTON	07/10/2018	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:37000-518-08	1 in 1 CARTON	07/01/2019	
2		8 in 1 BLISTER PACK; Type 0: Not a Combination		

2		Product		
3	NDC:37000-518-02	2 in 1 POUCH; Type 0: Not a Combination Product	01/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/10/2018	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 11/2023 The Procter & Gamble Manufacturing Company