VICKS DAYQUIL NYQUIL ULTRA CONCENTRATED COLD AND FLUacetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl

The Procter & Gamble Manufacturing Company

VICKS® DayQuil™ NyQuil™ ULTRA CONCENTRATED COLD & FLU, LiquiCaps™ Convenience Pack

VICKS® DayQuil™ ULTRA CONCENTRATED COLD & FLU, LiquiCaps™

Drug Facts

DayQuil™ Cold & Flu

Active Ingredient (in each LiquiCap)

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Phenylephrine HCl 5 mg

Purpose

Pain reliever/fever reducer

Cough suppressant

Nasal decongestant

Uses

temporarily relieves common cold/flu symptoms:

- nasal congestion
- cough due to minor throat & bronchial irritation
- sore throat
- headache
- minor aches & pains
- fever

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

- 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
- trouble urinating due to enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema

Ask a doctor or pharmacist before use if you are

taking the blood thinning drug warfarin

When using this product do not use more than directed

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

store at no greater than 25°C

Inactive ingredients

FD&C Yellow No. 5, FD&C Yellow No. 6, gelatin, glycerin, lecithin, mica, polyethylene glycol, polyvinyl acetate phthalate, povidone, sorbitol sorbitan solution, titanium dioxide, water

Questions?

1-800-362-1683

TAMPER EVIDENT: DO NOT USE IF PRINTED SEAL UNDER CAP IS MISSING OR DAMAGED.

DIST. BY PROCTER & GAMBLE,

CINCINNATI OH 45202

VICKS® NyQuil™ ULTRA CONCENTRATED COLD & FLU, LiquiCaps™

Drug Facts

NyQuil™ Cold & Flu

Active Ingredient (in each LiquiCap)

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Doxylamine succinate 6.25 mg

Purpose

Pain reliever/fever reducer

Cough suppressant

Antihistamine

Uses

temporarily relieves common cold/flu symptoms:

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

Warnings

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- 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

- 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, 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
- 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

store at no greater than 25°C

Inactive ingredients

D&C Yellow No. 10, FD&C Blue No. 1, gelatin, glycerin, lecithin, mica, polyethylene glycol, polyvinyl acetate phthalate, povidone, sorbitol sorbitan solution, titanium dioxide, water

Questions?

1-800-362-1683

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CINCINNATI OH 45202

PRINCIPAL DISPLAY PANEL

VICKS® DayQuil™ | VICKS® NyQuil™ ULTRA CONCENTRATED COLD & FLU, LiquiCaps™ MAX STRENGTH

2X VALUE PACK†

VICKS® DayQuil™

ULTRA CONCENTRATED

COLD & FLU

Acetaminophen, Phenylephrine HCl, Dextromethorphan HBr

Headache, Fever, Sore Throat, Minor Aches & Pains

Nasal Congestion, Sinus Pressure

Cough

25% SMALLER*

EASY TO SWALLOW

SWALLOW

Non-Drowsy

48 DAYQUIL LIQUICAPS™

VICKS® NyQuil™

ULTRA CONCENTRATED

COLD & FLU

Acetaminophen, Doxylamine Succinate, Dextromethorphan HBr

Headache, Fever, Sore Throat, Minor Aches & Pains

Sneezing, Runny Nose

Cough

25% SMALLER*

EASY TO SWALLOW

Nighttime Relief

48 DAYQUIL LIQUICAPS™ 48 NYQUIL LIQUICAPS™ 96 TOTAL LIQUICAPS™







VICKS DAYQUIL NYQUIL ULTRA CONCENTRATED COLD AND FLU

acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69423-802

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:69423-802- 96	1 in 1 CARTON; Type 0: Not a Combination Product	07/11/2023	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1		48
Part 2		48

Part 1 of 2

VICKS DAYQUIL ULTRA CONCENTRATED COLD AND FLU

acetaminophen, dextromethorphan hbr, phenylephrine hcl capsule, liquid filled

Product Information

Item Code (Source) NDC:69423-800

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		
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg		
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg		

Inactive Ingredients			
Ingredient Name	Strength		
MICA (UNII: V8A1AW0880)			
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
POVIDONE (UNII: FZ 989GH94E)			
WATER (UNII: 059QF0KO0R)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)			
GELATIN (UNII: 2G86QN327L)			
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)			
POLYVINYL ACETATE PHTHALATE (UNII: 58QVG85GW3)			
SORBITOL (UNII: 506T60A25R)			
GLYCERIN (UNII: PDC6A3C0OX)			

Product Characteristics			
Color	orange	Score	no score
Shape	OVAL	Size	15mm
Flavor		Imprint Code	DQ
Contains			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/11/2023	

Part 2 of 2

VICKS NYQUIL ULTRA CONCENTRATED COLD AND FLU

acetaminophen, dextromethorphan hbr, doxylamine succinate capsule, liquid filled

Product Information

Item Code (Source)	NDC:69423-801
Route of Administration	ORAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg		
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg		
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg		

Inactive Ingredients		
Ingredient Name	Strength	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POVIDONE (UNII: FZ989GH94E)		
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)		
SORBITOL (UNII: 506T60A25R)		
MICA (UNII: V8A1AW0880)		
GLYCERIN (UNII: PDC6A3C0OX)		
WATER (UNII: 059QF0KO0R)		
GELATIN (UNII: 2G86QN327L)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)		
POLYVINYL ACETATE PHTHALATE (UNII: 58QVG85GW3)		

Product Characteristics			
Color	green	Score	no score
Shape	OVAL	Size	15mm
Flavor		Imprint Code	NQ
Contains			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M012	07/11/2023		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M012	07/11/2023		

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 10/2023

The Procter & Gamble Manufacturing Company