VICKS DAYQUIL ULTRA CONCENTRATED COLD AND FLU- acetaminophen, dextromethorphan hbr, phenylephrine hcl capsule, liquid filled
The Procter & Gamble Manufacturing Company

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

Drug Facts

Active ingredients (in each LiquiCap)

Acetaminophen 325 mg
Dextromethorphan HBr 10 mg
Phenylephrine HCl 5 mg

Active ingredients (in each LiquiCap) 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 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

PRINCIPAL DISPLAY PANEL - 48 LiquiCaps™

MAX STRENGTH

100% MORE LIQUICAPS†

VICKS®

DayQuil™

ULTRA CONCENTRATED

COLD & FLU

Acetaminophen, Phenylephrine HCl, Dextromethorphan HBr

Headache, Fever, Sore Throat, Minor Aches & Pains

Nasal Congestion, Sinus Pressure

Cough

Non-Drowsy

25% SMALLER*

EASY TO SWALLOW

48 LIQUICAPS™

VICKS DAYQUIL ULTRA CONCENTRATED COLD AND FLU

acetaminophen, dextromethorphan hbr, phenylephrine hcl capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69423-800
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POLYVINYL ACETATE PHTHALATE (UNII: 58QVG85GW3)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
GLYCERIN (UNII: PDC6A3C0OX)	
POVIDONE (UNII: FZ989GH94E)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
MICA (UNII: V8A1AW0880)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
GELATIN (UNII: 2G86QN327L)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-800-48	1 in 1 CARTON	07/06/2023	
1		48 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC Monograph Drug	M012	07/06/2023	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 10/2024 The Procter & Gamble Manufacturing Company