VICKS DAYQUIL AND VICKS NYQUIL HIGH BLOOD PRESSURE COLD AND FLUacetaminophen, dextromethorphan hydrobromide, doxylamine succinate The Procter & Gamble Manufacturing Company

VICKS ® DayQuil™ NyQuil™ HIGH BLOOD PRESSURE COLD & FLU Liquid Convenience Pack

NyQuil HIGH BLOOD PRESSURE 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 & bronchial irritation
- sore throat
- headache
- minor aches & 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 (30 mL each) 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	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 28.23 mg
- store at no greater than 25°C and do not refrigerate

Inactive ingredients

citric acid, FD&C blue no. 1, FD&C red no. 40, flavor, glycerin, propylene glycol, saccharin sodium, sodium benzoate, sodium citrate, sorbitol, sucralose, water, xanthan gum

Questions?

1-800-362-1683

DIST. BY: PROCTER & GAMBLE,

CINCINNATI, OH 45202

DayQuil HIGH BLOOD PRESSURE Cold & Flu

Drug Facts

Active ingredients (in each 30 mL)

Acetaminophen 650 mg Dextromethorphan HBr 20 mg

Purpose

Pain reliever/Fever reducer Cough suppressant

Uses

temporarily relieves common cold/flu symptoms:

- 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 4 doses (30 mL each) 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
- 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.

Stop use and ask a doctor if

- pain or cough get worse or last 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 30 mL every 4 hrs children 4 to under 12 yrs ask a doctor children under 4 yrs do not use

Other information

- each 30 mL contains: sodium 26.06 mg
- store at no greater than 25°C and do not refrigerate

Inactive ingredients

citric acid, FD&C yellow no. 6, flavor, glycerin, propylene glycol, saccharin sodium, sodium benzoate, sodium citrate, sorbitol, sucralose, water, xanthan gum

Questions?

1-800-362-1683

TAMPER EVIDENT: Do not use if printed shrinkband seal around the neck is

broken or missing

DIST. BY: PROCTER & GAMBLE,

CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - Kit Carton

FAST RELIEF

DAY & NIGHT PACK

Vicks DayQuil and NyQuil HIGH BLOOD PRESSURE

Vicks DayQuil HIGH BLOOD PRESSURE

COLD & FLU

Acetaminophen, Dextromethorphan HBr

Headache, Fever, Sore Throat, Minor Aches & Pains

Cough

DECONGESTANT FREE

SPECIALLY FORMULATED

HBP

Vicks NyQuil HIGH BLOOD PRESSURE

COLD & FLU

Acetaminophen, Doxylamine Succinate, Dextromethorphan HBr

Headache, Fever, Sore Throat, Minor Aches & Pains

Sneezing, Runny Nose

Cough

DECONGESTANT FREE

2 BOTTLES - 1 DAYQUIL/1 NYQUIL; 8 FL OZ (236 mL) EACH; TOTAL 16 FL OZ (472 mL)



VICKS DAYQUIL AND VICKS NYQUIL HIGH BLOOD PRESSURE COLD AND FLU

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69423-974

I	Packaging			
4	# Item Code	Package Description	Marketing Start Date	Marketing End Date
]	NDC:69423-974- 16	1 in 1 PACKAGE; Type 0: Not a Combination Product	07/12/2022	
2	NDC:69423-974- 12	1 in 1 PACKAGE; Type 0: Not a Combination Product	07/09/2024	

Quant	Quantity of Parts			
Part #	Package Quantity	age Quantity Total Product Quantity		
Part 1	1 BOTTLE, PLASTIC	236 mL		
Part 2	1 BOTTLE, PLASTIC	236 mL		

Part 1 of 2

VICKS DAYQUIL HIGH BLOOD PRESSURE COLD AND FLU

acetaminophen, dextromethorphan liquid

Product Information		
Item Code (Source)	NDC:69423-972	
Route of Administration	ORAL	

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	650 mg in 30 mL	
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 30 mL	

Inactive Ingredients		
Ingredient Name	Strength	
XANTHAN GUM (UNII: TTV12P4NEE)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)		
GLYCERIN (UNII: PDC6A3C0OX)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SACCHARIN SODIUM (UNII: SB8ZUX40TY)		
SODIUM BENZOATE (UNII: O/245FE5EU)		

SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics			
Color	orange	Score	
Shape		Size	
Flavor	CITRUS	Imprint Code	
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423- 972-08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:69423- 972-06	177 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	07/12/2022		

Part 2 of 2

VICKS NYQUIL HIGH BLOOD PRESSURE COLD AND FLU

acetaminophen, dextromethorphan hydrobromide, and doxylamine succinate liquid

Product Information		
Item Code (Source)	NDC:69423-973	
Route of Administration	ORAL	

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	650 mg in 30 mL	
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg in 30 mL	
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	12.5 mg in 30 mL	

Inactive Ingredients	
Ingredient Name	Strength
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SORBITOL (UNII: 506T60A25R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
XANTHAN GUM (UNII: TTV12P4NEE)	
GLYCERIN (UNII: PDC6A3C0OX)	

Product Characteristics			
Color	red	Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

	Packaging			
7	# Item Code	Package Description	Marketing Start Date	Marketing End Date
:	NDC:69423- 973-08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:69423- 973-06	177 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	07/12/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/12/2022	

Labeler - The Procter & Gamble Manufacturing Company (004238200)