VICKS DAYQUIL SEVERE AND VICKS NYQUIL SEVERE COLD AND FLUacetaminophen, phenylephrine hydrochloride, dextromethorphan hydrobromide, guaifenesin and doxylamine succinate The Procter & Gamble Manufacturing Company

Vicks ® DayQuil™ Severe and Vicks ® NyQuil™ Severe Cold & Flu LiquiCaps™ Convenience Pack

DayQuil™ SEVERE Cold & Flu

Drug Facts

Active ingredients (in each LiquiCap)

Acetaminophen 325 mg
Dextromethorphan HBr 10 mg
Guaifenesin 200 mg
Phenylephrine HCl 5 mg

Purpose

Pain reliever/Fever reducer Cough suppressant Expectorant Nasal decongestant

Uses

temporarily relieves common cold/flu symptoms:

- nasal congestion
- sinus congestion & pressure
- cough due to minor throat & bronchial irritation
- minor aches & pains
- headache
- fever
- sore throat
- reduces swelling of nasal passages
- temporarily restores freer breathing through the nose
- promotes nasal and/or sinus drainage
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive.

Warnings

Liver warning:

This product contains acetaminophen.

Severe liver damage may occur if you take

- more than 8 LiquiCaps in 24 hrs, 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, chronic bronchitis, 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 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
- do not exceed 8 LiquiCaps per 24 hrs

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

Other information

do not exceed 25^oC

Inactive ingredients

ammonium hydroxide, FD&C Blue No. 1, FD&C Red No. 40, gelatin, glycerine, polyethylene glycol, povidone, propylene glycol, purified water, shellac, simethicone, sorbitol sorbitan solution, titanium dioxide

Questions? 1-800-362-1683 NyQuil™ SEVERE Cold & Flu 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 hrs, 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 hours

adults & children 12 2 LiquiCaps with water

yrs & over every 4 hrs

children 4 to under 12 yrs ask a doctor

children under 4 yrs do not use

Other information

do not exceed 25°C

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 - 12 Capsule Convenience Pack

MAX

STRENGTH

Vicks ® DayQuil™ Severe and Vicks ® NyQuil™ Severe Cold & Flu LiquiCaps™

VICKS®

DayQuil ™

SEVERE

COLD& FLU

Acetaminophen, Guaifenesin, Phenylephrine HCl, Dextromethorphan HBr

Headache, Fever, Sore Throat, Minor Aches & Pains

Chest Congestion, Thins & Loosens Mucus

Nasal Congestion & Sinus Pressure Cough

Non-Drowsy

8 DayQuil LiquiCaps

NyQuil ™

SEVERE

COLD& FLU

Acetaminophen, PhenylephrineHCl, Doxylamine Succinate, Dextromethorphan HBr

Headache, Fever, Sore Throat, Minor Aches & Pains

Nasal Congestion, Sinus Pressure

Sneezing, Runny Nose

Cough

Nighttime Relief

4 NyQuil LiquiCaps: TOTAL 12 LIQUICAPS



VICKS DAYQUIL SEVERE AND VICKS NYQUIL SEVERE COLD AND FLU

acetaminophen, phenylephrine hydrochloride, dextromethorphan hydrobromide, guaifenesin and doxylamine succinate kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:37000-934

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:37000-934-12	1 in 1 PACKAGE	07/04/2019		
2	NDC:37000-934-24	1 in 1 PACKAGE	07/04/2019		
3	NDC:37000-934-48	1 in 1 PACKAGE	07/04/2019		
4	NDC:37000-934-72	1 in 1 PACKAGE	07/04/2019		

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BLISTER PACK	8
Part 2	1 BLISTER PACK	4

Part 1 of 2

VICKS DAYQUIL SEVERE COLD AND FLU

acetaminophen, phenylephine hcl, dextromethorphan hydrobromide, and guaifenesin capsule, liquid filled

Product Information

Item Code (Source) NDC:37000-517

Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) **GUAIFENES IN** 200 mg ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII: 36209ITL9D) **ACETAMINOPHEN** 325 mg **DEXTROMETHORPHAN HYDROBROMIDE** (UNII: 9D2RTI9KYH) **DEXTROMETHORPHAN** 10 mg (DEXTROMETHORPHAN - UNII:7355X3ROTS) **HYDROBROMIDE** PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE -**PHENYLEPHRINE** 5 mg UNII:1WS297W6MV) **HYDROCHLORIDE**

Inactive Ingredients			
Ingredient Name	Strength		
AMMONIA (UNII: 5138Q19F1X)			
SHELLAC (UNII: 46N107B710)			
SORBITOL (UNII: 506T60A25R)			
FD&C RED NO. 40 (UNII: WZB9127XOA)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
GELATIN (UNII: 2G86QN327L)			
GLYCERIN (UNII: PDC6A3C0OX)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
POVIDONE (UNII: FZ 989GH94E)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
WATER (UNII: 059QF0KO0R)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			

Product Characteristics				
Color	orange Score ne		no score	
Shape	BULLET	Size	16mm	
Flavor		Imprint Code	DS	
Contains				

Pa	Packaging					
#	Item Package Description		Marketing Start Date	Marketing End Date		
1		1 in 1 CARTON				
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product				

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

Part 2 of 2

VICKS NYQUIL SEVERE COLD AND FLU

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

Product Information	
Item Code (Source)	NDC:37000-518
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: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg	
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg	
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg	

Inactive Ingredients			
Ingredient Name	Strength		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
GELATIN (UNII: 2G86QN327L)			
GLYCERIN (UNII: PDC6A3C0OX)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
POVIDONE (UNII: FZ989GH94E)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
WATER (UNII: 059QF0KO0R)			
SORBITOL (UNII: 506T60A25R)			

Product Characteristics			
Color	green Score n		no score
Shape	BULLET	Size	16mm
Flavor		Imprint Code	NS
Contains			

Pa	Packaging					
#	# Item Package Description		Marketing Start Date	Marketing End Date		
1		1 in 1 CARTON				
1		4 in 1 BLISTER PACK; Type 0: Not a Combination Product				

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

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/04/2019	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 10/2023 The Procter & Gamble Manufacturing Company