EQUATE MAXIMUM STRENGTH NIGHTTIME SEVERE COLD AND FLU SOFTGELS-acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hydrochloride capsule, liquid filled WALMART INC.

equateTM MAXIMUM STRENGTH NIGHTTIME SEVERE Cold & Flu SOFTGELS

Drug Facts

Active ingredients (in each softgel)

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 softgels 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, and 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,

Overdose warning: 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 softgels per 24 hrs

adults & children 12 yrs & over	2 softgels 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 controlled room temperature 20-25°C (68-77°F) • protect from light, heat, and moisture

Inactive ingredients

D&C Yellow No.10, FD&C Blue No.1, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, sorbitol sorbitan solution, titanium dioxide

Questions or comments?

1-888-287-1915

Compare to Vicks® NyQuil™ Severe Cold & Flu LiquiCaps™ active ingredients*

Relief of:

- Headache, fever, sore throat, minor aches and pains
- Nasal/sinus congestion and sinus pressure
- Sneezing, runny nose
- Cough

Satisfaction Guaranteed

For more information call 1-888-287-1915 or visit Walmart.com/help

DISTRIBUTED BY:

Walmart Inc.,

Bentonville, AR 72716

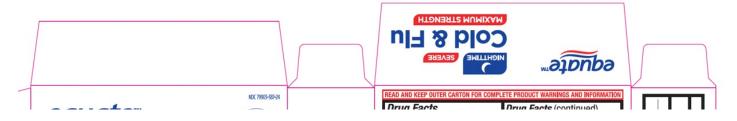
PRODUCT OF INDIA

*This product is not manufactured or distributed by The Procter & Gamble Company, owner of the registered trademarks Vicks[®] NyQuil™ LiquiCaps™.

READ AND KEEP OUTER CARTON FOR COMPLETE PRODUCT WARNINGS AND INFORMATION

TEMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OR TAMPERING.

Packaging





Walmart Inc., Bentonville, AR 72716 PRODUCT OF INDIA *This product is not



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Other information ■ store at controlls

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D&C Blue No.1, gelatin, glycerin

Questions or comments? 1-888-287-1915

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Inactive ingredients

D&C Yellow No.10, FD&C Blue No.1, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, sorbitol sorbitan solution, titanium dioxide

Questions or comments? 1-888-287-1915

EQUATE MAXIMUM STRENGTH NIGHTTIME SEVERE COLD AND FLU SOFTGELS

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

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79903-387
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
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		
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg		

Inactive Ingredients	
Ingredient Name	Strength
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SORBITOL (UNII: 506T60A25R)	
SORBITAN (UNII: 6092ICV9RU)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Prod	luct	Cha	arac	cter	istics

Color	areen	Score	no score

Shape	OVAL	Size	18mm
Flavor		Imprint Code	A07
Contains			

l	Packaging					
	# Item Code Package Description		Marketing Start Date	Marketing End Date		
	1 NDC:79903-387-24	2 in 1 CARTON	09/10/2025			
	1	12 in 1 BLISTER PACK; 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	09/10/2025		

Labeler - WALMART INC. (051957769)

Revised: 9/2025 WALMART INC.