

**DAYTIME NIGHTTIME COLD AND FLU RELIEF MULTI-SYMP TOM-
acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine
hcl**

P & L Development, LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug Facts

Active ingredients in Daytime (in each softgel)

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Phenylephrine HCL 5 mg

Active ingredients in Nighttime (in each softgel)

Acetaminophen 325 mg

Dextromethorphan HBr 15 mg

Doxylamine succinate 6.25 mg

Purpose for Daytime

Pain reliever/fever reducer

Cough suppressant

Nasal decongestant

Purpose for Nighttime

Pain reliever/fever reducer

Cough suppressant

Antihistamine

Uses

DAYTIME

- temporarily relieves common cold and flu symptoms
 - cough due to minor throat and bronchial irritation
 - nasal congestion
 - headache

- minor aches and pains
- fever
- sore throat

NIGHTTIME

- temporarily relieves common cold and flu symptoms
 - cough due to minor throat and bronchial irritation
 - sore throat
 - headache
 - minor aches and pain
 - fever
 - runny nose and sneezing

Warnings

DAYTIME and NIGHTTIME

Liver warning: These products contain acetaminophen. Severe liver damage may occur if you take:

- more than 4,000 mg of acetaminophen in 24 hours
- 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

DAYTIME and NIGHTTIME

- 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

DAYTIME

- liver disease
- heart disease

- diabetes
- thyroid disease
- high blood pressure
- cough that occurs with too much phlegm (mucus)
- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema

NIGHTTIME

- liver disease
- glaucoma
- cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts such as occur with smoking, asthma, chronic bronchitis or emphysema
- trouble urinating due to an enlarged prostate gland

Ask a doctor or pharmacist before use if you are

DAYTIME

- taking the blood thinning drug warfarin

NIGHTTIME

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers

When using this product,

DAYTIME

do not exceed recommended dosage.

NIGHTTIME

- **do not exceed recommended dosage**
- excitability may occur, especially in children
- avoid alcoholic drinks
- marked drowsiness may occur
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

DAYTIME

- nervousness, dizziness or sleeplessness occur
- pain, cough, and nasal congestion 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.

NIGHTTIME

- 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.

DAYTIME and NIGHTTIME

Overdose warning: Taking more than the recommended dose can cause liver damage. 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

DAYTIME

- **do not take more than directed (see Overdose warning)**
- do not take more than 4 doses in 24 hours
- swallow whole; do not crush, chew, or dissolve
- adults and children 12 years and over: take 2 softgels with water every 4 hours
- children under 12 years : do not use

Nighttime:

- **do not take more than directed (see Overdose warning)**
- do not take more than 4 doses in 24 hours
- swallow whole; do not crush, chew, or dissolve
- adults and children 12 years and over: take 2 softgels with water every 6 hours
- children under 12 years: do not use

Other information

DAYTIME and NIGHTTIME

- store between 15°-30°C (59°-86°F)
- avoid excessive heat

Inactive ingredients

Daytime butylated hydroxyanisole, butylated hydroxytoluene, FD&C yellow #6, gelatin, glycerin, mannitol, polyethylene glycol, povidone, propylene glycol, purified water, sorbitan, sorbitol, white ink

Nighttime D&C yellow #10, FD&C blue #1, gelatin, glycerin, mannitol, polyethylene glycol, povidone, propylene glycol, purified water, sorbitan, sorbitol, white ink

Questions or comments?

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

Principal Display Panel

Daytime

multi-symptom

cold & flu relief

acetaminophen (pain reliever / fever reducer)

dextromethorphan HBr (cough suppressant)

phenylephrine HCL (nasal decongestant)

- non-drowsy
- alcohol-free
- antihistamine-free

softgels**

(**liquid-filled capsules)

nighttime

multi-symptom

cold & flu relief

acetaminophen (pain reliever / fever reducer)

dextromethorphan HBr (cough suppressant)

doxylamine succinate (antihistamine)

- alcohol-free

(**liquid-filled capsules)

When using Daytime and Nighttime products, carefully read the labeling to ensure correct dosing.

†Compare to the active ingredients in Vicks® Dayquil® and Nyquil® Cold & Flu LiquiCaps®

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TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

DISTRIBUTED BY: PL Developments

200 Hicks Street, Westbury, NY 11590

Product Label



Daytime Cold & Flu Softgel

Drug Facts

Active ingredients (in each softgel)

- Acetaminophen 325 mg.....Pain reliever/fever reducer
- Dextromethorphan HBr 15 mg.....Cough suppressant
- Phenylephrine HCl 5 mg.....Nasal decongestant

Uses ■ temporarily relieves common cold and flu symptoms: ■ cough due to minor throat and bronchial irritation ■ sore throat ■ headache ■ minor aches and pains ■ fever ■ runny nose & sneezing

Warnings Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take: ■ more than 4,000 mg of acetaminophen in 24 hours ■ 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 redness ■ blisters ■ rash

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.

Nighttime Cold & Flu Softgel

Drug Facts

Active ingredients (in each softgel)

- Acetaminophen 325 mg.....Pain reliever/fever reducer
- Doxylamine succinate 6.25 mg.....Antihistamine

Uses ■ temporarily relieves common cold and flu symptoms: ■ cough due to minor throat and bronchial irritation ■ sore throat ■ headache ■ minor aches and pains ■ fever ■ runny nose & sneezing

Warnings Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take: ■ more than 4,000 mg of acetaminophen in 24 hours ■ 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 redness ■ blisters ■ rash

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.

Daytime Cold & Flu Softgel

Drug Facts (continued)

Ask a doctor before use if you have ■ liver disease ■ gauscoma ■ cough that occurs with too much phlegm (mucus) ■ a breathing problem or chronic cough that lasts such as occurs with smoking, asthma, chronic bronchitis, or emphysema ■ trouble urinating due to an enlarged prostate gland

Ask a doctor or pharmacist before use if you are ■ taking the blood thinning drug warfarin ■ taking sedatives or tranquilizers

When using this product ■ do not exceed recommended dosage ■ be careful when driving a motor vehicle or operating machinery ■ alcohol, sedatives, and tranquilizers may increase drowsiness ■ marked drowsiness may occur ■ avoid alcoholic drinks

Stop use and ask a doctor if ■ fever gets worse or lasts more than 7 days ■ pain or cough 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

Keep out of reach of children. Overdose warning: Taking more than the recommended dose can cause liver damage. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) 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

■ do not take more than directed (see Overdose warning) ■ do not take more than 4 doses in 24 hours ■ swallow whole; do not crush, chew, or dissolve ■ adults and children 12 years and over: take 2 softgels with water every 4 hours ■ children under 12 years: do not use

Nighttime Cold & Flu Softgel

Drug Facts (continued)

Ask a doctor before use if you have ■ liver disease ■ gauscoma ■ cough that occurs with too much phlegm (mucus) ■ a breathing problem or chronic cough that lasts such as occurs with smoking, asthma, chronic bronchitis, or emphysema ■ trouble urinating due to an enlarged prostate gland

Ask a doctor or pharmacist before use if you are ■ taking the blood thinning drug warfarin ■ taking sedatives or tranquilizers

When using this product ■ do not exceed recommended dosage ■ be careful when driving a motor vehicle or operating machinery ■ alcohol, sedatives, and tranquilizers may increase drowsiness ■ marked drowsiness may occur ■ avoid alcoholic drinks

Stop use and ask a doctor if ■ fever gets worse or lasts more than 7 days ■ pain or cough 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


Keep out of reach of children. Overdose warning: Taking more than the recommended dose can cause liver damage. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) 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


■ do not take more than directed (see Overdose warning) ■ do not take more than 4 doses in 24 hours ■ swallow whole; do not crush, chew, or dissolve ■ adults and children 12 years and over: take 2 softgels with water every 6 hours ■ children under 12 years: do not use

<p>dextromethorphan HBr (cough suppressant)</p> <p>phenylephrine HCl (nasal decongestant)</p> <ul style="list-style-type: none"> • non-drowsy • alcohol-free • antihistamine-free <p>32 softgels** (**liquid-filled capsules)</p>	<p>48 total softgels</p>	<p>dextromethorphan HBr (cough suppressant)</p> <p>doxylamine succinate (antihistamine)</p> <ul style="list-style-type: none"> • alcohol-free <p>16 softgels** (**liquid-filled capsules)</p>
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When using Daytime and Nighttime products, carefully read the labeling to ensure correct dosing.
 *Compare to the active ingredients in Vicks® DayQuil® and NyQuil® Cold & Flu LiquiCaps®



5 9726 47248 7



Actual Size

PLD-C5704
FD006975

Distributed by: **PL Developments**
200 Hicks Street, Westbury, NY 11590

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF BUSTLE UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

This product is not manufactured or distributed by The Procter & Gamble Company. Vicks®, DayQuil®, NyQuil®, and LiquiCaps® are registered trademarks of The Procter and Gamble Company.

<p>Daytime Cold & Flu Softgel</p> <p>Drug Facts (continued)</p> <p>Other Information</p> <ul style="list-style-type: none"> ■ store between 15-30°C (59-86°F) ■ avoid excessive heat <p>Inactive ingredients butylated hydroxyanisole, butylated hydroxytoluene, FD&C yellow #6, gelatin, glycerin, mannitol, polyethylene glycol, polyethylene glycol, purified water, sorbitan, sorbitol, white ink</p> <p>Questions or comments? Call 1-877-753-3935 Monday-Friday 9AM-5PM EST</p>	<p>Nighttime Cold & Flu Softgel</p> <p>Drug Facts (continued)</p> <p>Other Information</p> <ul style="list-style-type: none"> ■ store between 15-30°C (59-86°F) ■ avoid excessive heat <p>Inactive ingredients D&C yellow #10, FD&C blue #1, gelatin, glycerin, mannitol, polyethylene glycol, polyethylene glycol, purified water, sorbitan, sorbitol, white ink</p> <p>Questions or comments? Call 1-877-753-3935 Monday-Friday 9AM-5PM EST</p>
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WELLNESS BASICS Daytime Nighttime Multi-Symptom Cold and Flu Relief

DAYTIME NIGHTTIME COLD AND FLU RELIEF MULTI-SYMPATOM

acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59726-050
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59726-050-24	1 in 1 KIT; Type 0: Not a Combination Product	05/31/2019	12/27/2024

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
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Part 1	16 BLISTER PACK	16
Part 2	32 BLISTER PACK	32

Part 1 of 2

NIGHTTIME COLD AND FLU RELIEF

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate capsule

Product Information

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	15 mg
DOXYLAMINE SUCCINATE (UNII: V9B19B5Y12) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg

Inactive Ingredients

Ingredient Name	Strength
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
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)	
SORBITAN (UNII: 6O92ICV9RU)	
SORBITOL (UNII: 506T60A25R)	
MANNITOL (UNII: 3OWL53L36A)	

Product Characteristics

Color	green	Score	no score
Shape	CAPSULE	Size	20mm
Flavor		Imprint Code	P30
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	16 in 1 CARTON		
1	1 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	05/31/2019	12/27/2024

Part 2 of 2

DAYTIME COLD AND FLU RELIEF

acetaminophen, dextromethorphan hbr, phenylephrine hcl capsule

Product Information

Route of Administration ORAL

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
BUTYLATED HYDROXYANISOLE (UNII: REK4960K2U)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITAN (UNII: 6O92ICV9RU)	
SORBITOL (UNII: 506T60A25R)	
MANNITOL (UNII: 3OWL53L36A)	

Product Characteristics

Color	orange	Score	no score
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Shape	CAPSULE	Size	20mm
Flavor		Imprint Code	P19
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		32 in 1 CARTON		
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	05/31/2019	12/27/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	05/31/2019	12/27/2024

Labeler - P & L Development, LLC (800014821)

Revised: 3/2023

P & L Development, LLC