COLD AND FLU DAYTIME, NIGHTTIME, MAXIMUM STRENGTH- acetaminophen, dextromethorphan hbr, doxylamine succinate, guaifenesin, phenylephrine hcl Walgreen Company

Walgreens 44-640677-22

Active ingredients (in each caplet) (Daytime Severe Cold & Flu)

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

Purpose

Pain reliever/fever reducer Cough suppressant Expectorant Nasal decongestant

Active ingredients (in each caplet) (Nighttime Severe Cold & Flu)

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 and flu symptoms:
 - minor aches and pains
 - nasal congestion
 - headache
 - fever
 - sore throat
 - sinus congestion and pressure
 - cough due to minor throat and bronchial irritation
 - cough to help you sleep (Nighttime only)
 - runny nose and sneezing (Nighttime only)

- 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 (Daytime only)

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take

- with other drugs containing acetaminophen
- more than 4,000 mg of acetaminophen in 24 hours
- 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.
- if you have ever had an allergic reaction to this product or any of its ingredients

Ask a doctor before use if you have

- liver disease
- diabetes
- high blood pressure
- glaucoma (Nighttime only)
- thyroid disease
- heart disease
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema (Daytime only)
- difficulty in urination due to enlargement of the prostate gland
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma,

chronic bronchitis, or emphysema (Nighttime only)

Ask a doctor or pharmacist before use if you are

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers (Nighttime only)

When using this product

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

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- 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. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Do not take DAYTIME and NIGHTTIME products at the same time.

Directions

- do not take more than directed
- do not take more than 8 caplets of Daytime and Nighttime products in any 24-hour period
- adults and children 12 years and over: take 2 caplets with water every 4 hours
- children under 12 years: ask a doctor

Other information

- each caplet contains: sodium 3 mg
- TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive ingredients (Daytime only)

corn starch, crospovidone, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, polysorbate 80, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, talc, titanium dioxide

Inactive ingredients (Nighttime only)

black iron oxide, corn starch, crospovidone, D&C yellow #10 aluminum lake, FD&C blue #1 aluminum lake, FD&C yellow #6 aluminum lake, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, talc, titanium dioxide

Ouestions or comments?

1-800-426-9391

Principal Display Panel

DAY & NIGHT PACK

NDC 0363-6406-08

Walgreens

Compare to the active ingredients in Vicks® DayQuil® Severe Cold & Flu and Vicks® NyQuil® Severe Cold & Flu††

DAYTIME Severe Cold & Flu

ACETAMINOPHEN / PAIN RELIEVER /

FEVER REDUCER

DEXTROMETHORPHAN HBr / COUGH

SUPPRESSANT

GUAIFENESIN / EXPECTORANT PHENYLEPHRINE HCI / NASAL

DECONGESTANT

Maximum Strength

32 CAPLETS
ACTUAL SIZE

NIGHTTIME

Severe

Cold & Flu

ACETAMINOPHEN / PAIN RELIEVER /

FEVER REDUCER

DEXTROMETHORPHAN HBr / COUGH

SUPPRESSANT

DOXYLAMINE SUCCINATE / ANTIHISTAMINE

PHENYLEPHRINE HCI / NASAL

DECONGESTANT

Maximum Strength

16 CAPLETS ACTUAL SIZE

48 TOTAL CAPLETS

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

Do Not Take Daytime and

Nighttime Products at the Same Time.

PARENTS: Learn about teen medicine abuse www.StopMedicineAbuse.org

†Our pharmacists recommend the Walgreens brand. We invite you to compare to national brands.
††This product is not manufactured or distributed by The Procter & Gamble Company, owner of the registered trademark Vicks® DayQuil® Severe Cold & Flu and Vicks® NyQuil® Severe Cold & Flu.
50844 REV0722A64067722

DISTRIBUTED BY: WALGREEN CO.
200 WILMOT RD., DEERFIELD, IL 60015
Walgreens
100 % SATISFACTION
GUARANTEED
walgreens.com
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oorn starch, crospovidone, D&C yellow #10 aluminum lake, FD&C blue #1 aluminum lake, FD&C yellow #6 aluminum lake, magnesium Inactive ingredients (Nighttime only) black iron oxide,

auminum laks, magnesium stearate, matiodextrin, microcrystalline cellulose, polyethylans glycol, polysorbate 860, polyvinkyl alcohol, polycorbate 860, polyvinkyl alcohol, talc, listenia, silison dioxdes, sodium starch glycolate, stearic acid, talc, listenium dioxide. Inactive ingredients (Daytime only) corn starch, crospovidone, FD&C red #40 auminum lake, FD&C yellow #6

- see end flap for expiration date and lot number
- store at 25°C (7°F); excursions permitted between 15°-30°C (59°-86°F)
- Other information each caplet contains: sodium 3 mg
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Drug Facts (continued)

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- Milh any other drug containing acetaminopher, brescription or acetaminopher, ask a doctor or phermadst.

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- if you have ever had an allergic reaction to this product or any of its

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Mightlime Products at the

Do Not Take Daytime and

Sore throat warning: If sore throat is severe, pexsists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or more than a dod or promptly. ar nord 154 ni rendponimatsos lo gen 000.4, nentr stom ∎ nord stom of 2010 of

Dung Facts (continued)

Phenylephine HCl 5 mg..... Dextromethorphan HBr 10 mg...
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■ children under 12 years: ask a doctor Ask a doctor or pharmacist before use if you are adults and drildren 12 years and over. take 2 caplets with water every

Directions at the more than 8 caplets of Daylime and Mightime products

at on not take more than 8 caplets of Daylime and Mightime products
in any 24-hour pendo.

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Our pharmacists recommend the Walgreens brand.
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(duestions or comments? 1-800-426-9391

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Drug Facts (continued)

Do not take DYYIIME and WIGHTTIME products at the same time.

critical for adults as well as for children even if you do not notice any signs or symptoms. If pregnant or breast-feeding, sak a health protessional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control (senter right away, trompt medical alternion is contact a children are provided and the contact and con

■ cough comes back or occurs with resh or headache that lasts. These could be signs of a serious condition. ■ uew symptoms occur

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m nervousness, dizziness, or sleeplessness occur Stop use and ask a doctor if

(Nightime only)

a alcohol, sedatives, and tranquifizers may increase drowsiness
(Nightime only) ■ be careful when driving a motor vehicle or operating machinery.

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Drug Facts (continued)

LOT & EXP ONLY DO NOT PRINT TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

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ITEM 907684 W00000-0000

LÓO% SATISFACTION GUARANTEED

W30RG1121-F REV1122

DISTRIBUTED BY: WALGREEN CO. 200 WILMOT RD., DEERFIELD, IL 60015

■ with other drugs containing acetaminophen рашэде шэу оссиг и уои таке Liver warming: This product contains acetaminophen. Severe liver

■ rough to help you sleep (Nightlime only)
■ roundy accessing (Nightlime only)
■ reduces sand sneezing (Nightlime only)
■ reduces surelling of lessen be greated as members in the lost of the stores freet beginning a promotor measures access the companion of the second of the suns access resonance a promotor access to the suns a promotor access to the suns access to the suns access a promotor access to the suns access to the su

Drug Facts (continued)

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> Active ingredients (in each caplet) (Nighttime Severe Cold & Flu) Purpose Dextormethorphan HBr 10 mg Cough suppressant Custormethorphan HBr 10 mg Expectorant Custlenesin 200 mg Expectorant Massal decongestant mg Respectorant Custormethorphine HCl 5 mg Cough specifications of the cough specific management of the cough specific management

(Daytime Severe Cold & Flu) Active ingredients (in each caplet)

Drug Facts

COMPLETE PRODUCT INFORMATION

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NDC 0363-6406-22

DAY & NIGHT PACK

Walgreens

Severe Cold & Flu

ACETAMINOPHEN / PAIN RELIEVER / FEVER REDUCER
DEXTROMETHORPHAN HBr / COUGH SUPPRESSANT GUAIFENESIN / EXPECTURANT Phenylephrine hct / Nasal Decongestant

Maximum Strength

32 CAPLETS

NIGHTTIME

Severe Cold & Flu

ACETAMINOPHEN / PAIN RELIEVER / FEVER REDUCER DOXYLAMINE SUCCINATE / ANTIHISTAMINE PHENYLEPHRINE HCI / NASAL DECONGESTANT

Maximum Strength

ACTUAL SIZE

16 CAPLETS

ACTUAL SIZE

48 TOTAL CAPLETS

Walgreens 44-640677-22

B-2201-640677-22HR REV0722A64067722

COLD AND FLU DAYTIME, NIGHTTIME, MAXIMUM STRENGTH

acetaminophen, dextromethorphan hbr, doxylamine succinate, guaifenesin, phenylephrine hcl kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0363-6406

Pac	kag	ing

# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:0363- 6406-22	1 in 1 PACKAGE, COMBINATION; Type 0: Not a Combination Product	08/01/2015	

Quantity of Parts

4	, 6. 7 6. 6.		
Part #	Package Quantity	Total Product Quantity	
Part 1	4 BLISTER PACK	32	
Part 2	2 BLISTER PACK	16	

Part 1 of 2

COLD AND FLU DAYTIME, MAXIMUM STRENGTH

acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl tablet, film coated

Product Information

Route of Administration ORAL

Active Ingredient/Active Moiety

Active ingredient/Active Molety		
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
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients	
Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
CROSPOVIDONE, UNSPECIFIED (UNII: 2S7830E561)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	orange	Score	no score
Shape	OVAL	Size	19mm
Flavor		Imprint Code	44;640
Contains			

Packaging					
	# Item Package Description			Marketing Start Date	Marketing End Date
	1		8 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 Drug	M012	02/27/2014	

Part 2 of 2

COLD AND FLU NIGHTTIME, MAXIMUM STRENGTH

acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl tablet, film coated

Product Information

Route of Administration ORAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg
DEVELOPMENT OF THE PROPERTY OF	DEVEDOMETHORDHAM	

DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9DZKTI9KTH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	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
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
STARCH, CORN (UNII: O8232NY3SJ)	
CROSPOVIDONE, UNSPECIFIED (UNII: 2S7830E561)	
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)	
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYSORBATE 80 (UNII: 60ZP39ZG8H)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	green	Score	no score
Shape	OVAL	Size	19mm
Flavor		Imprint Code	44;677
Contains			

Packaging					
# Item Package Description		Marketing Start Date	Marketing End Date		
1		8 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 Drug	M012	08/01/2015			

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	08/01/2015	

Labeler - Walgreen Company (008965063)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(0363-6406) , pack(0363-6406)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(0363-6406)

Establishment				
Name	Address	ID/FEI	Business Operations	
LNK International, Inc.		868734088	manufacture(0363-6406)	

Establishment				
Name	Address	ID/FEI	Business Operations	
LNK International, Inc.		117025878	manufacture(0363-6406)	

Revised: 12/2023 Walgreen Company