

**SINUS PRESSURE AND PAIN DAYTIME NIGHTTIME MAXIMUM STRENGTH-
acetaminophen, diphenhydramine hcl, guaifenesin, phenylephrine hcl
Walgreen Company**

Walgreens 44-615694-09-SMH

Active ingredients (in each caplet) (daytime)

Acetaminophen 325 mg
Guaifenesin 200 mg
Phenylephrine HCl 5 mg

Purpose

Pain reliever
Expectorant
Nasal decongestant

Active ingredients (in each caplet) (nighttime)

Acetaminophen 325 mg
Diphenhydramine HCl 12.5 mg
Phenylephrine HCl 5 mg

Purpose

Pain reliever
Antihistamine/cough suppressant
Nasal decongestant

Uses

- temporarily relieves these common cold symptoms:
 - nasal congestion
 - headache
 - minor aches and pains
 - sinus congestion and pressure
 - cough (***nighttime only***)
 - runny nose and sneezing (***nighttime only***)
- temporarily 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

- 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.

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.
- with any other product containing diphenhydramine, even one used on skin **(nighttime only)**

Ask a doctor before use if you have

- heart disease
- high blood pressure
- liver disease
- thyroid disease
- diabetes
- difficulty in urination due to enlargement of the prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- cough that occurs with too much phlegm (mucus)
- a breathing problem such as emphysema or chronic bronchitis **(nighttime only)**
- glaucoma **(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)**
- alcohol, sedatives, and tranquilizers may increase drowsiness **(nighttime only)**
- use caution when driving a motor vehicle or operating machinery **(nighttime only)**
- avoid alcoholic beverages **(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
- new symptoms occur
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- 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 12 caplets in any 24-hour period
- adults and children 12 years and over: take 2 caplets every 4 hours
- children under 12 years: do not use

Other information

- **each caplet contains:** sodium 3 mg (***daytime only***)
- **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, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, talc, titanium dioxide

Inactive ingredients (nighttime only)

corn starch, croscarmellose sodium, crospovidone, FD&C blue #1 aluminum lake, FD&C blue #2 aluminum lake, iron oxide yellow, magnesium stearate, methacrylic acid and ethyl acrylate copolymer, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, sodium bicarbonate, stearic acid, talc, titanium dioxide

Questions or comments?

1-800-426-9391

Principal display panel

DAY & NIGHT PACK

NDC 0363-6156-09

Walgreens

• **WALGREENS** •
PHARMACIST RECOMMENDED†

DAYTIME	NIGHTTIME
Sinus Pressure & Pain	Sinus Pressure & Pain
ACETAMINOPHEN	ACETAMINOPHEN
PAIN RELIEVER	PAIN RELIEVER
GUAIFENESIN	DIPHENHYDRAMINE HCl
EXPECTORANT	ANTIHISTAMINE / COUGH SUPPRESSANT
PHENYLEPHRINE HCl	PHENYLEPHRINE HCl
NASAL DECONGESTANT	NASAL DECONGESTANT
Maximum Strength	Maximum Strength
12 CAPLETS	8 CAPLETS
ACTUAL SIZE	ACTUAL SIZE

TOTAL 20 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.**

†Our pharmacists recommend the Walgreens brand.
We invite you to compare to national brands.

50844 REV0824B61569409

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DEERFIELD, IL 60015

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No print/No varnish area

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

Our pharmacists recommend the Walgreens brand. We invite you to compare to national brands.
50844 REV0824B61569409

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

<p>Drug Facts</p> <p>Active ingredients (in each caplet) (daytime)</p> <ul style="list-style-type: none"> Acetaminophen 325 mg Guafenesin 200 mg Phenylephrine HCl 5 mg 	<p>Purpose</p> <p>Pain reliever</p>
<p>Active ingredients (in each caplet) (nighttime)</p> <ul style="list-style-type: none"> Diphenhydramine HCl 12.5 mg Phenylephrine HCl 5 mg 	<p>Purpose</p> <p>Pain reliever</p>

Uses

- temporarily relieve these common cold symptoms:
 - nasal congestion and pressure
 - cough (nighttime only)
 - runny nose and sneezing (nighttime only)
- temporarily soothes nasal and/or throat irritation
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passages of both mucus and make coughs more productive (daytime only)

Warnings

Liver warnings: This product contains acetaminophen. Severe liver damage may occur if you take more than 4,000 mg of acetaminophen in 24 hours.

■ even one dose containing acetaminophen more than 4,000 mg may cause liver damage.

■ Allergic alert: Acetaminophen may cause an allergic reaction. Symptoms may include:

- hives
- itching
- swelling of the face, lips, tongue, or throat
- difficulty breathing
- skin rash

■ If you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, Parkinson's disease, 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.

■ Do not use with any other product containing diphenhydramine, even one used on skin (nighttime only).

Do not use

- with any other drug containing acetaminophen (prescription or over-the-counter), if you are on a drug containing acetaminophen, ask a doctor or pharmacist.
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, Parkinson's disease, 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 are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, Parkinson's disease, 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.

PEEL HERE FOR MORE DRUG FACTS

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NDC 0363-6156-09

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DAY & NIGHT PACK

Walgreens 

DAYTIME

Sinus Pressure & Pain

ACETAMINOPHEN

PAIN RELIEVER
GUAIFENESIN
EXPECTORANT
PHENYLEPHRINE HCl
NASAL DECONGESTANT

Maximum Strength

12 CAPLETS  ACTUAL SIZE

NIGHTTIME

Sinus Pressure & Pain

ACETAMINOPHEN

PAIN RELIEVER
DIPHENHYDRAMINE HCl
ANTIHISTAMINE / COUGH SUPPRESSANT
PHENYLEPHRINE HCl
NASAL DECONGESTANT

Maximum Strength

8 CAPLETS  ACTUAL SIZE

TOTAL 20 CAPLETS

Do Not Take Daytime and Nighttime Products at the Same Time.

B-2201-61569409HRV
REV0824B61569409

ADHESIVE AREA

Drug Facts (continued)

Ask a doctor before use if you have

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

Drug Facts (continued)

- use caution when driving a motor vehicle or operating machinery (nighttime only)
- avoid alcoholic beverages (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
- new symptoms occur
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- 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.

ADHESIVE AREA

ADHESIVE AREA

Drug Facts (continued)

Directions

- do not take more than directed
- do not take more than 12 caplets in any 24-hour period
- adults and children 12 years and over: take 2 caplets every 4 hours
- children under 12 years: do not use

Other information

- each caplet contains: sodium 3 mg (daytime only)
- 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, croscopolone, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, talc, titanium dioxide

Drug Facts (continued)

Inactive ingredients (nighttime only)

corn starch, croscarmellose sodium, croscopolone, FD&C blue #1 aluminum lake, FD&C blue #2 aluminum lake, iron oxide yellow, magnesium stearate, methacrylic acid and ethyl acrylate copolymer, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, sodium bicarbonate, stearic acid, talc, titanium dioxide

Questions or comments? 1-800-426-9391

Walgreens Pharmacist Recommended. Our pharmacists recommend the Walgreens brand. We invite you to compare to national brands.

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ITEM 886729
W00000-0000-0


PAPER BOX MULTI-LAYER TRAY
how2recycle.info
W30RG0423-F REV-0125

ADHESIVE AREA

Walgreens 44-615694

SINUS PRESSURE AND PAIN DAYTIME NIGHTTIME MAXIMUM STRENGTH

acetaminophen, diphenhydramine hcl, guaifenesin, phenylephrine hcl kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-6156
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-6156-09	1 in 1 CARTON; Type 0: Not a Combination Product	06/02/2018	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BLISTER PACK	12
Part 2	1 BLISTER PACK	8

Part 1 of 2

SINUS PRESSURE AND PAIN DAYTIME MAXIMUM STRENGTH

acetaminophen, guaifenesin, phenylephrine hcl tablet, film coated

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
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)	
CROSPVIDONE, UNSPECIFIED (UNII: 2S7830E561)	
FD&C RED NO. 40 ALUMINUM LAKE (UNII: 6T47AS764T)	
FD&C YELLOW NO. 6 ALUMINUM LAKE (UNII: GYP6Z2JR6Q)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
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;615
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		12 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	06/02/2018	

Part 2 of 2

SINUS PRESSURE AND PAIN NIGHTTIME MAXIMUM STRENGTH

acetaminophen, diphenhydramine hcl, phenylephrine hcl tablet, film coated

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
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	12.5 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
CROSPVIDONE, UNSPECIFIED (UNII: 2S7830E561)	
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	
FD&C BLUE NO. 2 ALUMINUM LAKE (UNII: 4AQJ3LG584)	

FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHACRYLIC ACID AND ETHYL ACRYLATE COPOLYMER (UNII: NX76LV5T8J)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	blue	Score	no score
Shape	OVAL	Size	19mm
Flavor		Imprint Code	44;694
Contains			

Packaging

#	Item Code	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	06/02/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	06/02/2018	

Labeler - Walgreen Company (008965063)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(0363-6156)

Establishment

Name	Address	ID/FEI	Business Operations
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LNK International, Inc.		832867837	manufacture(0363-6156) , pack(0363-6156)
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Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(0363-6156)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(0363-6156)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(0363-6156)

Revised: 12/2025

Walgreen Company