

**DAY AND NIGHT SINUS RELIEF MAXIMUM STRENGTH- acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, guaifenesin, phenylephrine hydrochloride  
Wal-Mart Stores, Inc.**

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**Maximum Strength Day & Night Sinus Relief**

***Active ingredients (in each Softgel)***

**Sinus Daytime**

**Acetaminophen 325 mg**

Dextromethorphan HBr 10 mg

Guaifenesin 200 mg

Phenylephrine HCl 5 mg

**Sinus Nighttime**

**Acetaminophen 325 mg**

Dextromethorphan HBr 10 mg

Doxylamine succinate 6.25 mg

Phenylephrine HCl 5 mg

***Purposes***

**Sinus Daytime**

**Pain reliever**

Cough suppressant

Expectorant

Nasal decongestant

**Sinus Nighttime**

**Pain reliever**

Cough suppressant

Antihistamine

Nasal decongestant

***Uses***

- temporarily relieves:
- nasal congestion
- headache
- cough
- minor aches and pains
- sinus congestion and pressure
- 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 12 softgels in 24 hours, which is the maximum daily amount
- 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.

## **Ask a doctor before use if you have**

- liver disease
- heart disease
- diabetes
- high blood pressure
- thyroid disease
- trouble urinating due to an enlarged prostate gland
- glaucoma (**Nighttime only**)
- a breathing problem such as emphysema or chronic bronchitis (**Nighttime only**)
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- cough that occurs with too much phlegm (mucus)

## **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 use more than directed**
- excitability may occur, especially in children (**Nighttime only**)
- marked drowsiness may occur (**Nighttime only**)
- alcohol, sedatives, and tranquilizers may increase drowsiness (**Nighttime only**)

- avoid alcoholic drinks (**Nighttime only**)
- be careful when driving a motor vehicle or operating machinery (**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.**

**Overdose warning:** Taking more than the recommended dose (overdose) may 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**

- **do not take more than directed (see Overdose warning)**
- do not take more than 12 softgels in any 24-hour period
- adults and children 12 years of age and older: take 2 softgels every 4 hours
- children under 12 years of age: do not use

### **Other information**

- store at room temperature 15°-30°C (59°-86°F) and avoid excessive heat.

### **Inactive ingredients**

**(Daytime only)** FD&C yellow #6, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, purified water, sorbitol sorbitan solution and white ink

**(Nighttime only)** D&C yellow #10, FD&C blue #1, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, purified water, sorbitol sorbitan solution and white ink

### **Questions or comments?**

Call toll free: **1-888-287-1915**

### **PRINCIPAL DISPLAY PANEL - Carton Label**

Maximum Strength Day and Night Sinus Relief Combo 24ct

**DAY & NIGHT TWIN PACK**



**MAXIMUM STRENGTH**  
Daytime  
**Sinus Relief**  
ACETAMINOPHEN 325mg  
Dextromethorphan HBr 10mg  
Guaifenesin 200mg  
Phenylephrine HCl 5mg

**MAXIMUM STRENGTH**  
Nighttime  
**Sinus Relief**  
ACETAMINOPHEN 325mg  
Dextromethorphan HBr 10mg  
Doxylamine succinate 4.25mg  
Phenylephrine HCl 5mg

**DAY & NIGHT TWIN PACK**


NDC 0085-16104

**Compare to Mucinex® Sinus-Max® Day and Night Maximum Strength Active Ingredients\***

Maximum Strength\*\*

Fast Dissolving Softgels

## Sinus Relief

Day



**ACETAMINOPHEN**  
Pain Reliever

**Dextromethorphan HBr**  
Cough Suppressant

**Guaifenesin**  
Expectorant

**Phenylephrine HCl**  
Nasal Decongestant

- Relieves sinus pressure, headache & congestion
- Controls cough
- Thins & loosens mucus



16 SOFTGELS  
(Liquid Filled Capsules)

Night



**ACETAMINOPHEN**  
Pain Reliever

**Dextromethorphan HBr**  
Cough Suppressant

**Doxylamine succinate**  
Antihistamine

**Phenylephrine HCl**  
Nasal Decongestant

- Relieves nasal congestion, sinus pressure & pain
- Controls cough
- Relieves runny nose & sneezing



8 SOFTGELS  
(Liquid Filled Capsules)

24 TOTAL

AGES 12+

LOT NO:

EXP DATE:

COATING  
FREE AREA



6 81131 27526 2

**PARENTS:**  
Learn about teen medicine abuse.  
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RECYCLED FIBER MULTI-LAYER BOX TRAY  
Printed on 100% Recycled Paper

\*Compare to Mucinex Sinus-Max Day and Night Maximum Strength Active Ingredients

\*\*Per 4-hour dose

Do not take the Daytime and Nighttime softgels at the same time. Do not take more than a total of 12 softgels in a 24-hour period.

Do not take the first dose of the Nighttime softgels sooner than 4 hours after the last dose of the Daytime softgels unless directed by a doctor.

Take only as directed.

Keep cartons for complete product information.

**Caution:** Do not use if carton is damaged or a blister pack on blister is broken or missing.

**Warnings:** Do not use if you have liver disease, heart disease, diabetes, high blood pressure, thyroid disease, trouble urinating due to an enlarged prostate gland, glaucoma, or breathing problems such as emphysema or chronic bronchitis.

**Directions:** Do not take more than directed. Do not take more than 2 tablets in any 24-hour period. Do not take more than 16 tablets in any 28-day period. Do not take more than 8 tablets in any 24-hour period. Do not take more than 2 tablets every 4 hours. Do not use if you are pregnant or breastfeeding.

**Other Information:** Store at room temperature (59°-86°F) and avoid excessive heat.

**Questions or comments?** 1-800-347-0915

**Drug Facts (continued)**

**Warnings:** Liver warnings: This product contains acetaminophen. Severe liver damage may occur if you take: more than 12 tablets in 24 hours, which is the maximum daily amount; with other drugs containing acetaminophen; or more alcoholic drinks daily while using this product. Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include: skin redness, blisters, rash. If a skin reaction occurs, stop use and seek medical help right away.

**Directions:** Do not take more than directed (see **Overdose warning**). Do not take more than 2 tablets in any 24-hour period. Do not use if you are pregnant or breastfeeding. Do not take more than 2 tablets every 4 hours. Do not use if you are under 12 years of age; do not use.

**Other Information:** Store at room temperature (59°-86°F) and avoid excessive heat.

**Inactive ingredients:** (Daytime only) FD&C yellow #1, pectin, glycerin, polyethylene glycol, polystyrene glycol, purified water, sorbitol sorbitan solution, and white ink. (Nighttime only) FD&C yellow #10, FD&C blue #1, glycerin, glycerin, polyethylene glycol, polystyrene glycol, purified water, sorbitol sorbitan solution, and white ink.

**Drug Facts**

**Active ingredients (in each Softgel) Purpose:**  
 **sinus Daytime**  
 Acetaminophen 325 mg... Pain reliever  
 Dextromethorphan HBr 10 mg... Cough suppressant  
 Guaifenesin 200 mg... Expectorant  
 Phenylephrine HCl 5 mg... Nasal decongestant

**Active ingredients (in each Softgel) Purpose:**  
 **sinus Nighttime**  
 Acetaminophen 325 mg... Pain reliever  
 Dextromethorphan HBr 10 mg... Cough suppressant  
 Doxylamine succinate 6.25 mg... Antihistamine  
 Phenylephrine HCl 5 mg... Nasal decongestant

**Uses:** temporarily relieves: nasal congestion, headache, cough, minor aches and pains, sinus congestion and pressure, runny nose and sneezing (Nighttime only). temporarily promotes nasal and/or sinus drainage. helps loosen phlegm (mucus) and thin bronchial secretions to clear the bronchial passageways of bothersome 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 12 tablets in 24 hours, which is the maximum daily amount; with other drugs containing acetaminophen; or more alcoholic drinks daily while using this product. Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include: skin redness, blisters, rash. If a skin reaction occurs, stop use and seek medical help right away.

**Directions:** Do not take more than directed (see **Overdose warning**). Do not take more than 2 tablets in any 24-hour period. Do not take more than 16 tablets in any 28-day period. Do not take more than 8 tablets in any 24-hour period. Do not take more than 2 tablets every 4 hours. Do not use if you are pregnant or breastfeeding.

**Other Information:** Store at room temperature (59°-86°F) and avoid excessive heat.

**Inactive ingredients:** (Daytime only) FD&C yellow #1, pectin, glycerin, polyethylene glycol, polystyrene glycol, purified water, sorbitol sorbitan solution, and white ink. (Nighttime only) FD&C yellow #10, FD&C blue #1, glycerin, glycerin, polyethylene glycol, polystyrene glycol, purified water, sorbitol sorbitan solution, and white ink.

## DAY AND NIGHT SINUS RELIEF MAXIMUM STRENGTH

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

### Product Information

**Product Type** HUMAN OTC DRUG **Item Code (Source)** NDC:49035-587

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-587-04	1 in 1 CARTON; Type 0: Not a Combination Product	01/31/2018	

### Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	2 BLISTER PACK	16
Part 2	1 BLISTER PACK	8

### Part 1 of 2

### DAY SINUS RELIEF

acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride capsule, liquid filled

## Product Information

**Route of Administration** ORAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

## Inactive Ingredients

Ingredient Name	Strength
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	
<b>GELATIN</b> (UNII: 2G86QN327L)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POVIDONE</b> (UNII: FZ989GH94E)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>SORBITAN</b> (UNII: 6O92ICV9RU)	
<b>WATER</b> (UNII: 059QF0KO0R)	

## Product Characteristics

<b>Color</b>	orange (clear)	<b>Score</b>	no score
<b>Shape</b>	CAPSULE (Oblong)	<b>Size</b>	25mm
<b>Flavor</b>		<b>Imprint Code</b>	PC26
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		2 in 1 CARTON		
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	01/31/2018	

## Part 2 of 2

### NIGHT SINUS RELIEF

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

#### Product Information

Route of Administration ORAL

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
<b>DOXYLAMINE SUCCINATE</b> (UNII: V9B19B5Y12) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1VS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

#### Inactive Ingredients

Ingredient Name	Strength
<b>POVIDONE</b> (UNII: FZ989GH94E)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>SORBITAN</b> (UNII: 6O92ICV9RU)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>D&amp;C YELLOW NO. 10</b> (UNII: 35SW5USQ3G)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>GELATIN</b> (UNII: 2G86QN327L)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	

#### Product Characteristics

<b>Color</b>	green (clear)	<b>Score</b>	no score
<b>Shape</b>	CAPSULE (Oblong)	<b>Size</b>	21mm
<b>Flavor</b>		<b>Imprint Code</b>	PC22
<b>Contains</b>			

#### Packaging

#	Item Code	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		
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### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/31/2018	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/31/2018	

**Labeler** - Wal-Mart Stores, Inc. (051957769)

### Establishment

Name	Address	ID/FEI	Business Operations
Humanwell PuraCap Pharmaceuticals (Wuhan) Co., Ltd		421293287	manufacture(49035-587)

Revised: 11/2024

Wal-Mart Stores, Inc.