

**SINUS PRESSURE AND PAIN MAXIMUM STRENGTH- acetaminophen,  
phenylephrine hcl tablet  
DOLGENCORP, LLC**

-----  
**Dollar General 44-502 RR**

***Active ingredients (in each caplet)***

Acetaminophen 325 mg  
Phenylephrine HCl 5 mg

***Purpose***

Pain reliever/fever reducer  
Nasal decongestant

***Uses***

- temporarily relieves these symptoms associated with hay fever or other upper respiratory allergies, and the common cold:
  - minor aches and pains
  - nasal congestion
  - headache
  - sinus congestion and pressure
- promotes sinus drainage
- temporarily reduces fever

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

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

- diabetes
- liver disease
- heart disease
- thyroid disease
- high blood pressure
- difficulty in urination due to enlargement of the prostate gland

**Ask a doctor or pharmacist before use if you are**

taking the blood thinning drug warfarin.

**When using this product**

**do not exceed recommended dosage.**

**Stop use and ask a doctor if**

- nervousness, dizziness, or sleeplessness occur
- pain or 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

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 accidental 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.

***Directions***

- **do not use more than directed**
- adults and children 12 years and over
  - take 2 caplets every 4 hours
  - do not take more than 10 caplets in 24 hours
- children under 12 years: ask a doctor

***Other information***

- **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°C-30°C (59°F-86°F)
- see end flap for expiration date and lot number

### ***Inactive ingredients***

corn starch, croscarmellose sodium, crospovidone, FD&C yellow #6 aluminum lake, magnesium stearate, microcrystalline cellulose, povidone, silicon dioxide, stearic acid

### ***Questions or comments?***

**1-888-309-9030**

### ***Principal display panel***

**DG™ | health**

**Compare to active ingredients of Sudafed PE® Sinus Pressure + Pain\***

**Maximum Strength**

**Sinus Pressure & Pain**

**Acetaminophen** • Pain Reliever/Fever Reducer  
**Phenylephrine HCl** • Nasal Decongestant

**Relieves:** • Sinus pressure & congestion  
• Sinus headache

Pseudoephedrine Free

Non Drowsy

**24 Caplets**

Actual Caplet Size

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

\*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark SUDAFED PE® SINUS PRESSURE + PAIN.

50844                      REV0820L50208

DISTRIBUTED BY OLD EAST MAIN CO.  
100 MISSION RIDGE  
GOODLETTSVILLE, TN 37072

**100% Satisfaction Guaranteed!**  
(888)309-9030

# Maximum Strength Sinus Pressure & Pain

**DG™** | health

Compare to active ingredients of **SUDAFED PE® SINUS PRESSURE + PAIN™**

# Maximum Strength Sinus Pressure & Pain

**Acetaminophen • Pain Reliever/Fever Reducer**  
**Phenylephrine HCl • Nasal Decongestant**

Relieves: • Sinus pressure & congestion  
• Sinus headache

**Pseudoephedrine Free**

Non Drowsy



**24 Caplets**

Actual Caplet Size



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

**NO PRINT/NO VARNISH LOT & EXP DATE**

\*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark SUDAFED PE® SINUS PRESSURE + PAIN.  
50844 REV0820L50208  
DISTRIBUTED BY OLD EAST MAIN CO.  
100 MISSION RIDGE  
GOODLETTSVILLE, TN 37072

**100% Satisfaction Guaranteed!**  
(888) 309-9030

**Drug Facts (continued)**  
**Other information**  
 ■ 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

**Drug Facts (continued)**  
**Inactive ingredients** corn starch, croscarmellose sodium, croscopolone, FD&C yellow #6 aluminum lake, magnesium stearate, microcrystalline cellulose, povidone, silicon dioxide, stearic acid

**Questions or comments?** 1-888-309-9030  
B-0315-502-08RR  
REV0820L50208

**Drug Facts**  
**KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION**

**Active ingredients (in each caplet)**  
 Acetaminophen 325 mg.....Pain reliever/fever reducer  
 Phenylephrine HCl 5 mg.....Nasal decongestant

**Purpose**  
 ■ temporarily relieves these symptoms associated with hay fever or other upper respiratory allergies, and the common cold:  
 ■ minor aches and pains ■ nasal congestion ■ headache  
 ■ sinus congestion and pressure  
 ■ promotes sinus drainage ■ temporarily reduces fever

**Uses**  
 ■ temporarily relieves these symptoms associated with hay fever or other upper respiratory allergies, and the common cold:  
 ■ minor aches and pains ■ nasal congestion ■ headache  
 ■ sinus congestion and pressure  
 ■ promotes sinus drainage ■ temporarily reduces fever

**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 2 weeks after stopping the MAOI drug, if you do not know if your prescription drug contains an MAOI, ask a doctor

**Directions**  
 ■ do not use more than directed  
 ■ adults and children 12 years and over  
 ■ take 2 caplets every 4 hours  
 ■ do not take more than 10 caplets in 24 hours  
 ■ children under 12 years: ask a doctor

**Stop use and ask a doctor if**  
 ■ nervousness, dizziness, or sleeplessness occur  
 ■ pain or 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  
 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 accidental 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.

**When using this product do not exceed recommended dosage.**

**Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin.**

**Ask a doctor before use if you have**  
 ■ diabetes ■ liver disease ■ heart disease ■ thyroid disease ■ high blood pressure  
 ■ difficulty in urination due to enlargement of the prostate gland

**Ingredients**  
 ■ if you have ever had an allergic reaction to this product or any of its ingredients

**Drug Facts (continued)**

**DG Health 44-502**

# SINUS PRESSURE AND PAIN MAXIMUM STRENGTH

acetaminophen, phenylephrine hcl tablet

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:55910-925
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

## Inactive Ingredients

Ingredient Name	Strength
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>CROSPVIDONE, UNSPECIFIED</b> (UNII: 2S7830E561)	
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	

## Product Characteristics

<b>Color</b>	orange	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	18mm
<b>Flavor</b>		<b>Imprint Code</b>	44;502
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-925-08	2 in 1 CARTON	06/23/2005	
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	06/23/2005	
--------------------	------	------------	--

**Labeler -** DOLGENCORP, LLC (068331990)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		832867837	manufacture(55910-925) , pack(55910-925)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		117025878	manufacture(55910-925)

Revised: 6/2024

DOLGENCORP, LLC