# SUDAFED PE SINUS PRESSURE PLUS PAIN- acetaminophen and phenylephrine hydrochloride tablet, film coated Kenvue Brands LLC

-----

#### Sudafed PE Sinus Pressure Plus Pain

# Drug Facts

Active ingredients (in each tablet)	Purpose
Acetaminophen 325 mg	Pain reliever/fever
Acetaminophen 525 mg	reducer
Phenylephrine HCl 5 mg	Nasal decongestant

#### Uses

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

# Warnings

#### Liver warning

This product contains acetaminophen. The maximum daily dose of this product is 10 tablets (3,250 mg acetaminophen) in 24 hours. 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

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged 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 dose

# 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.

# **Overdose warning**

In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222) 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 use more than directed (see overdose warning)

adults and children 12 years and over	<ul> <li>take 2 tablets every 4 hours</li> <li>do not take more than tablets in 24 hours</li> </ul>	10
--	---	----

children under 12 years ask a doctor

# Other information

store between 20-25°C (68-77°F)

do not use if blister unit is torn or broken

#### Inactive ingredients

carnauba wax, FD&C yellow no. 6 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, modified starch, polyethylene glycol, polysorbate 80, powdered cellulose, pregelatinized starch, sodium starch glycolate, titanium dioxide

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

call 1-888-217-2117 (toll-free) or 215-273-8755 (collect)

# PRINCIPAL DISPLAY PANEL

PREVIOUSLY SUDAFED PE <sup>®</sup> PRESSURE + PAIN NDC 50580-435-01

SUDAFED PE®

SINUS PRESSURE + PAIN

Acetaminophen, Phenylephrine HCl Pain Reliever/Fever Reducer, Nasal Decongestant

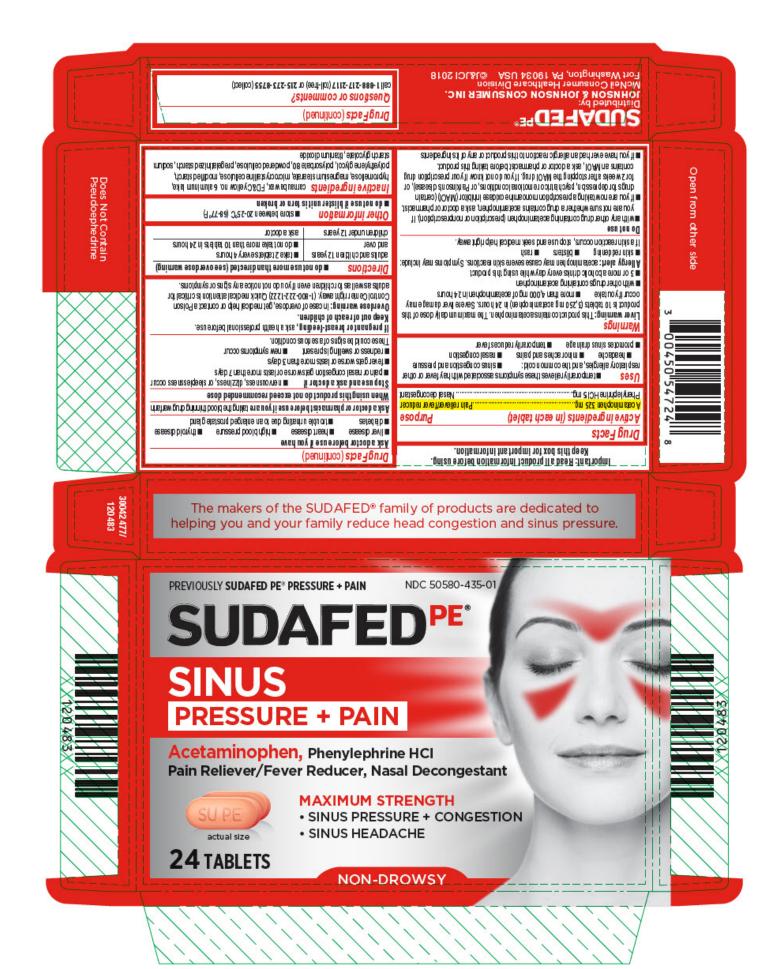
actual size

MAXIMUM STRENGTH

- SINUS PRESSURE + CONGESTION
- SINUS HEADACHE

24 TABLETS

NON-DROWSY



		PRESSURE PL			ed			
Product Info	rmation							
Product Type		HUMAN OTC DRUG	ltem Co	Code (Source) ND			IDC:50580-435	
Route of Admin	istration	ORAL						
Active Ingred	lient/Active	Moiety						
	Ingre	dient Name			Basis of St	rength	Strengt	
ACETAMINOPHEN	I (UNII: 36209ITI	9D) (ACETAMINOPHEN - U	JNII:362O9IT	L9D)	ACETAMINOPHEN		325 mg	
PHENYLEPHRINE JNII:1WS297W6MV)	HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - PHENYLEPHRINE HYDROCHLORIDE				5 mg			
Inactive Ingra	edients							
j.		Ingredient Name	9			S	trength	
CARNAUBA WAX	(UNII: R12CBM0E	-					<b>J</b>	
TITANIUM DIOXID								
FD&C YELLOW N	<b>0.6</b> (UNII: H77V	(EI93A8)						
ALUMINUM OXIDE	(UNII: LMI2606	933)						
HYPROMELLOSE,	UNSPECIFIED	(UNII: 3NXW29V3WO)						
MAGNESIUM STE	ARATE (UNII: 70	097M6I30)						
MICROCRYSTALL	INE CELLULOS	E (UNII: OP1R32D61U)						
POLYETHYLENE C	GLYCOL, UNSPI	ECIFIED (UNII: 3WJQ0SDV	V1A)					
POLYSORBATE 80	<b>0</b> (UNII: 60ZP39)	ZG8H)						
POWDERED CELL	ULOSE (UNII: S	MD1X3XO9M)						
SODIUM STARCH	GLYCOLATE T	YPE A (UNII: H8AV0SQX4)	D)					
Product Char	acteristics							
Color	orange (P	EACH)	Score no so		io score	score		
Shape	OVAL		Size 1		8mm			
Flavor			Imprint Code S		UPE;WL89			
Contains								
Packaging								
# Item Code	Ра	ckage Description		Mar	keting Start Date		ting End ate	
<b>1</b> NDC:50580- 435-01	2 in 1 CARTON		(	06/17/2019				
1	12 in 1 BLISTE Product	R PACK; Type 0: Not a Co	Combination					
Maxizatina	1 <b></b>	ion						
Marketing	intormat	ion						
Marketing	Applica	tion Number or Mon	ograph	Mai	rketing Start	Marke	eting End	

Category	Citation	Date	Date
OTC Monograph Drug	M012	06/17/2019	
	11012	00/1//2015	

# Labeler - Kenvue Brands LLC (118772437)

Revised: 11/2024

Kenvue Brands LLC