

SINUS PRESSURE PLUS PAIN PE- acetaminophen and phenylephrine hydrochloride tablet, coated
MARC GLASSMAN, INC.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

MAR-1120B-2022-1005

Drug Facts

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

Uses

- temporarily relieves these symptoms associated with hay fever or other upper 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 caplets (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 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.

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 take more than directed (see overdose warning)**

adults and children 12 years and over	<ul style="list-style-type: none"> ▪ take 2 caplets every 4 hours ▪ do not take more than 10 caplets in 24 hours
children	

CHILDREN
under 12
years

- ask a doctor

Other information

- store between 20-25°C (68-77°F) in a dry place
- retain carton for complete product information and warnings

Inactive ingredients

colloidal silicon dioxide, corn starch, croscarmellose sodium, crospovidone, FD&C yellow #6 aluminum lake, iron oxide yellow, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, pregelatinized starch, stearic acid, talc, titanium dioxide

PRINCIPAL DISPLAY PANEL

Marc's®

NDC 68998-220-06

†Compare to the active ingredients in Sudafed PE® Sinus Pressure + Pain

Maximum Strength, Non-Drowsy

Sinus Pressure + Pain PE

Acetaminophen, Phenylephrine HCl

PAIN RELIEVER/FEVER REDUCER, NASAL DECONGESTANT

For Relief of:

- Sinus Pressure + Congestion
- Sinus Headache

Actual Size

24 CAPLETS

Drug Facts (continued)

Other Information

- store between 20-25°C (68-77°F) in a dry place
- retain carton for complete product information and warnings

Inactive ingredients

colloidal silicon dioxide, corn starch, croscarmellose sodium, crospovidone,

Drug Facts (continued)

FD&C yellow #6 aluminum lake, iron oxide yellow, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, pregelatinized starch, stearic acid, talc, titanium dioxide

Drug Facts (continued)

Active ingredients (in each caplet)

Purpose

Pain reliever/fever reducer
 Acetaminophen 325 mg
 Phenylephrine HCl 5 mg
 Nasal decongestant

Uses

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

Warnings

This product contains acetaminophen. The maximum daily dose of this product is 10 caplets (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. Other drugs containing acetaminophen or 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.

Directions

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

Do not take more than directed (see overdose warning)

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.

Keep out of reach of children.

If pregnant or breast-feeding, ask a health professional before use. These could be signs of a serious condition.

- redness or swelling is present
- new symptoms occur
- fever gets worse or lasts more than 3 days
- pain or nasal congestion gets worse or lasts more than 7 days
- nervousness, dizziness, or sleeplessness occur

Stop use and ask a doctor if

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

Drug Facts (continued)

■ if you have ever had an allergic reaction to this product or any of its ingredients
 ■ liver disease
 ■ heart disease
 ■ thyroid disease
 ■ high blood pressure
 ■ trouble urinating due to an enlarged prostate gland



DO NOT USE IF BLISTER UNITS ARE TORN OR BROKEN

This product is not manufactured or distributed by McKel Consumer Healthcare, distributor of Sudafed PE® Sinus Pressure + Pain.

Distributed by:
 Marc Glassman, Inc.
 West 130th Street
 Cleveland, OH 44130



Maximum Strength • Non-Drowsy
Sinus Pressure + Pain PE



Maximum Strength • Non-Drowsy
Sinus Pressure + Pain PE

Acetaminophen, Phenylephrine HCl

PAIN RELIEVER/FEVER REDUCER, NASAL DECONGESTANT



24 CAPLETS

For Relief of:
 • Sinus Pressure + Congestion
 • Sinus Headache

NC

NC

NC

SINUS PRESSURE PLUS PAIN PE

acetaminophen and phenylephrine hydrochloride tablet, coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68998-220
Route of Administration	ORAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
CROSPVIDONE, UNSPECIFIED (UNII: 2S7830E561)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	orange	Score	no score
Shape	OVAL	Size	17mm
Flavor		Imprint Code	AAA;1120
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68998-220-06	2 in 1 CARTON	06/22/2020	
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 final	part341	06/22/2020	

Labeler - MARC GLASSMAN, INC. (094487477)

Revised: 10/2022

MARC GLASSMAN, INC.