# SINUS SEVERE- acetaminophen, guaifenesin, and phenylephrine hydrochloride tablet, coated CHAIN DRUG CONSORTIUM

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.

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#### 1119-PRV-2020-0823

#### Drug Facts

Active ingredients (in each caplet)	Purpose
Acetaminophen 325 mg	Pain reliever/fever
Acetaninophen 525 mg	reducer
Guaifenesin 200 mg	Expectorant
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
  - nasal congestion
  - minor aches and pains
- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive
- 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
- 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

# When using this product do not exceed recommended dosage

## 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

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	take 2 caplets every 4 hours swallow whole – do not crush, chew, or dissolve		
years and over	<ul> <li>do not take more than 10 caplets in 24 hours</li> </ul>		
children under 12 years	<ul> <li>ask a doctor</li> </ul>		

# Other information

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

#### **Inactive ingredients**

acesulfame potassium, colloidal silicon dioxide, croscarmellose sodium, crospovidone, flavor, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, pregelatinized starch, propylene glycol, stearic acid, talc, titanium dioxide

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

1-844-705-4384

## PRINCIPAL DISPLAY PANEL

Premier Value® COMPARE TO THE ACTIVE INGREDIENTS IN TYLENOL® SINUS SEVERE FOR ADULTS Sinus Severe DAYTIME • NON-DROWSY Acetaminophen, Phenylephrine HCl, Guaifenesin Pain Reliever / Fever Reducer, Nasal Decongestant, Expectorant Relieves: • Sinus Headache • Sinus Pressure Nasal Congestion •Mucus & Chest Congestion 24 COOL TASTE CAPLETS

ACTUAL SIZE



SINUS SEVERE

Product Information	ı					
Product Type	HU	MAN OTC DRUG	Item Code (S	em Code (Source) NDC:680		6-504
Route of Administration	OR	AL				
Active Ingredient/A	ctive Moiety					
	Ingredie	ent Name		Basis of	Strength	Strengtl
ACETAMINO PHEN (UNII:	0		362O9ITL9D)	ACETAMINOPH	-	325 mg
GUAIFENESIN (UNII: 495)	W7451VQ) (GUAI	FENESIN - UNII:495W745	51VQ)	GUAIFENESIN		200 mg
PHENYLEPHRINE HYDRO UNII:1WS297W6MV)	ENYLEPHRINE HYDRO CHLO RIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - PHENYLEPHRINE				5 mg	
Inactive Ingredients						
		Ingredient Name				Strength
ACESULFAME POTASSI	U <b>M</b> (UNII: 230V7	0				0
SILICON DIOXIDE (UNII:						
CROSCARMELLOSE SO		OL1HH48)				
CROSPOVIDONE (UNII: 2		,				
MAGNESIUM STEARATE		30)				
MALTODEXTRIN (UNII: 7		,				
CELLULOSE, MICROCR		II: OP1R32D6 1U)				
POLYETHYLENE GLYCO						
POLYVINYL ALCOHOL	(UNII: 532B59J99	90)				
PO VIDO NE, UNSPECIFIE	<b>D</b> (UNII: FZ989G	H94E)				
STARCH, PREGELATINIZ	ZED CORN (UNII	: O8232NY3SJ)				
PROPYLENE GLYCOL (U	JNII: 6 DC 9 Q 16 7 V	73)				
STEARIC ACID (UNII: 4EL	V7Z65AP)					
TALC (UNII: 7SEV7J4R1U)	1					
TITANIUM DIO XIDE (UN	II: 15FIX9V2JP)					
Product Characteris	stics					
Color	white	Score	Score		no score	
Shape	OVAL	Size	Size		19 mm	
Flavor	MINT	Imprint Cod	Imprint Code		AAA;1119	
Contains						
Packaging						
# Item Code	Pac	Package Description Ma		rketing Start Da	ate Marketi	ng End Dat
<b>1</b> NDC:68016-504-24 2 in	1 CARTON		11/0	1/2009		
			ation Product			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part341	11/01/2009				
OTC monograph final		11/01/2009				

# Labeler - CHAIN DRUG CONSORTIUM (101668460)

Revised: 9/2020

CHAIN DRUG CONSORTIUM