#### SINUS PAIN AND CONGESTION- acetaminophen, chlorpheniramine maleate, phenylephrine hcl CVS Pharmacy

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#### SINUS PAIN AND CONGESTION

#### Active ingredients (in each caplet) (Daytime Sinus Congestion & Pain)

Acetaminophen 325 mg Phenylephrine HCl 5 mg

#### Purpose

Pain reliever/fever reducer Nasal decongestant

#### Active ingredients (in each caplet) (Nighttime Sinus Congestion & Pain)

Acetaminophen 325 mg Chlorpheniramine maleate 2 mg Phenylephrine HCl 5 mg

#### Purpose

Pain reliever/fever reducer Antihistamine Nasal decongestant

#### Uses

- temporarily relieves these symptoms associated with hay fever or other respiratory allergies, and the common cold:
  - nasal congestion
  - headache
  - minor aches and pains
  - sinus congestion and pressure
  - runny nose and sneezing (Nighttime only)
- promotes sinus drainage
- helps clear nasal passages
- helps decongest sinus openings and passages
- 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

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

- heart disease
- high blood pressure
- liver disease
- thyroid disease
- diabetes
- difficulty in urination due to enlargement of the prostate gland
- a breathing problem such as emphysema or chronic bronchitis (*Nighttime only*)
- glaucoma (Nighttime only)

## 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 exceed recommended dosage
- use caution when driving a motor vehicle or operating machinery (*Nighttime only*)
- drowsiness may occur (Nighttime only)
- excitability may occur, especially in children (Nighttime only)
- alcohol, sedatives, and tranquilizers may increase drowsiness (Nighttime only)
- avoid alcoholic beverages (Nighttime only)

## 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.

# Read each section carefully. Do not take DAYTIME and NIGHTTIME products at the same time.

#### Directions

- do not take more than directed
- adults and children 12 years and over
  - take 2 caplets every 4 hours
  - swallow whole do not crush, chew, or dissolve
  - 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°-30°C (59°-86°F)
- see end flap for expiration date and lot number

# Inactive ingredients (Daytime only)

corn starch, crospovidone, D&C yellow #10 aluminum lake, FD&C blue #1 aluminum lake, FD&C red #40 aluminum lake, flavor, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, sucralose, talc, titanium dioxide

# Inactive ingredients (Nighttime only)

corn starch, crospovidone, flavor, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, sucralose, talc, titanium dioxide

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

1-800-426-9391

#### **Principal Display Panel**

♥CVSHealth<sub>®</sub>

Sinus

Actual With cool blast flavor Size Actual Size	DAYTIME - Non-Drowsy Sinus Pain & Congestion ACETAMINOPHEN Pain reliever, Fever reducer PHENYLEPHRINE HCI Nasal decongestant Relieves: Headache, Sinus pressure & Nasal congestion 12 CAPLETS With cool blast flavor Actual Size	Actual
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## **24 TOTAL CAPLETS**

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

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V-19849

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50844 REV0721C45546608

Do not take the **Daytime** and **Nighttime** caplets at the same time.



CVS 44-455C466

## SINUS PAIN AND CONGESTION

	nation						
Product Type	HUMAN	OTC DRUG	Item Co	de (Souro	e)	NDC:59779	-779
Packaging				Mark	eting Start	Marko	ting End
# Item Code	Pao	ckage Descrip	tion	Mark	Date		ate
<b>1</b> NDC:59779-779-	1 in 1 CARTON Product	N; Type 0: Not a C	ombination	07/26/20	05		
Quantity of Pai	rts						
Part #	Package C	Quantity		Tot	al Product Q	uantity	
Part 1 1 BLISTER PA	ACK		12				
Part 2 1 BLISTER P	ACK		12				
<b>Product Inform</b> Route of Adminis <sup>:</sup>		ORAL					
Active Ingredie	nt/Active	Moiety					
Active Ingredie		Moiety dient Name			Basis of S	itrength	Strengt
ACETAMINOPHEN (L	<b>Ingre</b> JNII: 36209ITL	<b>dient Name</b> L9D) (ACETAMINOP			ACETAMINOPHE	EN	<b>Strengt</b> 325 mg
ACETAMINOPHEN (U PHENYLEPHRINE HY	<b>Ingre</b> JNII: 36209ITL	<b>dient Name</b> L9D) (ACETAMINOP				EN E	
ACETAMINOPHEN (U PHENYLEPHRINE HY UNII:1W5297W6MV)	Ingre JNII: 36209ITL DROCHLORI	<b>dient Name</b> L9D) (ACETAMINOP			ACETAMINOPHE PHENYLEPHRIN	EN E	325 mg
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ACETAMINOPHEN (U PHENYLEPHRINE HY JNII:1WS297W6MV)	Ingre JNII: 36209ITL DROCHLORI Iients I: 08232NY3S	dient Name L9D) (ACETAMINOP IDE (UNII: 04JA59T IDE Ingredient	NSJ) (PHENYLE t Name		ACETAMINOPHE PHENYLEPHRIN	E E DE	325 mg 5 mg
ACETAMINOPHEN (UPHENYLEPHRINE HY UNII:1W5297W6MV)	Ingre JNII: 36209ITL DROCHLORI lients I: 08232NY3S NSPECIFIED	dient Name _9D) (ACETAMINOP IDE (UNII: 04JA59T IDE (UNII: 04JA59T Ingredient	NSJ) (PHENYLE <b>t Name</b> 1)		ACETAMINOPHE PHENYLEPHRIN	E E DE	325 mg 5 mg
ACETAMINOPHEN (U PHENYLEPHRINE HY JNII: 1WS 297W6MV) Inactive Ingred STARCH, CORN (UNII CROSPOVIDONE, UNI D&C YELLOW NO. 1	Ingre JNII: 36209ITL (DROCHLORI IEENTS I: 08232NY3S NSPECIFIED	dient Name (Job (ACETAMINOP (UNII: 04JA59T (UNII: 04JA59T (UNII: 257830E56: (UNII: 257830E56: (UNII: CQ:	NSJ) (PHENYLE <b>t Name</b> 1) 3XH3DET6)		ACETAMINOPHE PHENYLEPHRIN	E E DE	325 mg 5 mg
ACETAMINOPHEN (U PHENYLEPHRINE HY UNII: 1W5 297W6MV) Inactive Ingred STARCH, CORN (UNII CROSPOVIDONE, UN D&C YELLOW NO. 1 FD&C BLUE NO. 1 A	Ingre JNII: 36209ITL (DROCHLORI IEENTS I: 08232NY3S NSPECIFIED LO ALUMINUM LA	dient Name _9D) (ACETAMINOP IDE (UNII: 04JA59T Ingredient 5J) (UNII: 2S7830E56: M LAKE (UNII: CQ: AKE (UNII: J9EQA3	NSJ) (PHENYLE <b>t Name</b> 1) 3XH3DET6)		ACETAMINOPHE PHENYLEPHRIN	E E DE	325 mg 5 mg
ACETAMINOPHEN (U PHENYLEPHRINE HY UNII: 1W5 297W6MV) Inactive Ingred STARCH, CORN (UNII CROSPOVIDONE, UN D&C YELLOW NO. 1 FD&C BLUE NO. 1 A FD&C RED NO. 40 (I	Ingre JNII: 36209ITL (DROCHLORI IEINTS I: 08232NY3S NSPECIFIED LO ALUMINUM ALUMINUM LA UNII: WZ B912	dient Name L9D) (ACETAMINOP IDE (UNII: 04JA59T IDE (UNII: 04JA59T (UNII: 2S7830E56: M LAKE (UNII: CQ: AKE (UNII: J9EQA3 7XOA)	NSJ) (PHENYLE <b>t Name</b> 1) 3XH3DET6)		ACETAMINOPHE PHENYLEPHRIN	E E DE	325 mg 5 mg
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ΡΟν	POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)								
SILIC	SILICON DIOXIDE (UNII: ETJ7Z6XBU4)								
SOD	IUM STAR	RCH GLYC	OLATE T	YPE A POTATO (UN	NII: 5856J3G2A2	2)			
STE/	STEARIC ACID (UNII: 4ELV7Z65AP)								
SUC	RALOSE (	UNII: 96K6	UQ3ZD4)						
TALC	<b>C</b> (UNII: 7S	EV7J4R1U)	)						
TITA	NIUM DIO	XIDE (UN	III: 15FIX9\	/2JP)					
Pro	oduct Cł	naracte	eristics						
Colo	or		gree	n	Score			no sco	ore
Sha	ре		OVAL	-	Size			17mm	
Flav	/or		MENT	THOL	Imprint Cod	le		44;466	5
Con	tains								
Pac	ckaging								
#	ltem Code		Pack	age Descriptio	n	Mark	eting Start Date	Mar	keting End Date
1			LISTER PA	ACK; Type 0: Not a C	Combination				
		Product							
Marketing Information									
Ma	arketir	ng Inte	ormat	ion					
	Marketin	ng		tion Number or	Monograph	Ма	rketing Start	Ма	rketing End
	Marketir Categor	ng ry	Applica		Monograph		Date	Ma	rketing End Date
	Marketin	ng ry		tion Number or	Monograph	<b>Ma</b> 07/26	Date	Ма	
	Marketir Categor	ng ry	Applica	tion Number or	Monograph		Date	Ma	
отс	Marketin Categor Monograp	ng 'Y h Drug M	Applica	tion Number or	Monograph		Date	Ma	
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inactive in	ngredie	nts						
		Ingr	edient Name		Strength			
STARCH, CORN (UNII: 08232NY3SJ)								
CROSPOVIDONE, UNSPECIFIED (UNII: 2S7830E561)								
MAGNESIUM STEARATE (UNII: 70097M6I30)								
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)								
OLYETHYLE	NE GLYC	OL, UNSPECIFIED (	UNII: 3WJQ0SDW1A)					
POLYVINYL A	LCOHOL,	UNSPECIFIED (UNI	l: 532B59J990)					
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)								
SILICON DIO	KIDE (UNII	: ETJ7Z6XBU4)						
SODIUM STA	RCH GLYC	OLATE TYPE A PO	<b>TATO</b> (UNII: 5856J3G2A2	)				
TEARIC ACI	<b>D</b> (UNII: 4E	LV7Z65AP)						
SUCRALOSE	(UNII: 96K6	5UQ3ZD4)						
TALC (UNII: 79	-							
TITANIUM DI	OXIDE (UN	III: 15FIX9V2JP)						
	_							
Product C	haracte	eristics						
Color		white	Score		no score			
Shape		OVAL	Size	17mm				
lavor		MENTHOL	Imprint Cod	<b>e</b> 44;455				
Contains								
Packaging	J							
" Item		Deckere De		Marketing Start	Marketing End			
Code		Package De	scription	Date	Date			
1	12 in 1 E	BLISTER PACK; Type	0: Not a Combination					
	Product							
		_						
Marketi	ng Inf	ormation						
	-	ormation	nber or Monograph	Marketing Start	Marketing End			
Marketi Marketi Catego	ng	Application Nur	nber or Monograph tation	Marketing Start Date	Marketing End Date			
Marketi Catego	ng ry	Application Nur	<b>.</b> .		<b>-</b>			
Marketi	ng ry	Application Nur Ci	<b>.</b> .	Date	<b>-</b>			
Marketi Catego	ng ry	Application Nur Ci	<b>.</b> .	Date				
Marketi Catego OTC Monograp	ng ry oh Drug N	Application Nur Ci 1012	<b>.</b> .	Date				
Marketi Catego DTC Monograp <b>Marketi</b>	ng ry oh Drug M ng Inf	Application Nur Ci 1012 Ormation	tation	<b>Date</b> 06/28/2005	Date			
Marketi Catego DTC Monograp <b>Marketi</b> Marketi	ng ry oh Drug M ng Inf ng	Application Nur Ci 1012 Ormation Application Nur	<b>.</b> .	Date	<b>-</b>			
Marketi Catego DTC Monograp <b>Marketi</b>	ng ry oh Drug M ng Inf ng ry	Application Nur Ci 1012 Ormation Application Nur	tation <b>Contract</b>	Date 06/28/2005 Marketing Start	Date Marketing End			

Labeler - CVS Pharmacy (062312574)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
LNK International, Inc.		038154464	pack(59779-779)

Establishment				
Name	Address	ID/FE	3	<b>Business Operations</b>
LNK International, Inc.		83286783	7 manufactu	re(59779-779) , pack(59779-779)
Establishment				
Name	Ad	dress	ID/FEI	<b>Business Operations</b>
LNK International, Inc.			832867894	manufacture(59779-779)
Establishment				
Name	Ad	dress	ID/FEI	<b>Business Operations</b>
LNK International, Inc.			967626305	pack(59779-779)
Establishment				
Namo	٨d	drocs	ID/EEI	Business Operations

Name	Address	ID/FEI	<b>Business Operations</b>
LNK International, Inc.		117025878	manufacture(59779-779)

Revised: 1/2024

**CVS** Pharmacy