DAYTIME SINUS NIGHTTIME SINUS MAXIMUM STRENGTH- acetaminophen, diphenhydramine hcl, guaifenesin, phenylephrine hcl Target Corporation

Target 44-615694-SM

Active ingredients (in each caplet) (Sinus Day)

Acetaminophen 325 mg Guaifenesin 200 mg Phenylephrine HCl 5 mg

Purpose

Pain reliever Expectorant Nasal decongestant

Active ingredients (in each caplet) (Sinus Night)

Acetaminophen 325 mg Diphenhydramine HCl 12.5 mg Phenylephrine HCl 5 mg

Purpose

Pain reliever Antihistamine/cough suppressant Nasal decongestant

Uses

- temporarily relieves:
 - nasal congestion
 - headache
 - minor aches and pains
 - sinus congestion and pressure
 - cough (Nighttime only)
 - runny nose and sneezing (Nighttime only)
- temporarily promotes nasal and/or sinus drainage
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive (Daytime only)

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
- with any other product containing diphenhydramine, even one used on skin (Nighttime only)

Ask a doctor before use if you have

- heart disease
- liver disease
- diabetes
- thyroid disease
- high blood pressure
- difficulty in urination due to enlargement of the prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- cough that occurs with too much phlegm (mucus)
- 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
- excitability may occur, especially in children (Nighttime only)
- marked drowsiness may occur (Nighttime only)
- alcohol, sedatives, and tranquilizers may increase drowsiness (Nighttime only)
- avoid alcoholic beverages (Nighttime only)

• be careful when driving a motor vehicle or operating machinery (Nighttime only)

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.

In case of accidental overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

If taking NIGHTTIME and DAYTIME products, carefully read each section to ensure correct dosing. Do not take DAY & NIGHT at the same time.

Directions

- do not use more than directed
- do not take more than 12 caplets in any 24-hour period
- adults and children 12 years and over: take 2 caplets every 4 hours
- children under 12 years: do not use

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, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, talc, titanium dioxide

Inactive ingredients (Nighttime only)

corn starch, croscarmellose sodium, crospovidone, FD&C blue #1 aluminum lake, FD&C blue #2 aluminum lake, iron oxide yellow, magnesium stearate, methacrylic acid and

ethyl acrylate copolymer, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, sodium bicarbonate, stearic acid, talc, titanium dioxide

Questions?

Call 1-800-910-6874

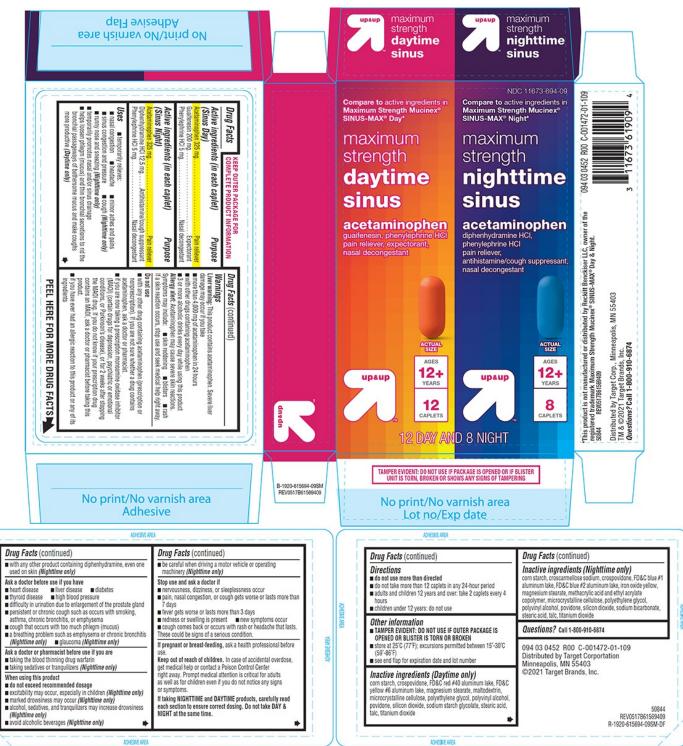
Principal Display Panel

	NDC 11673-694-09
	Compare to active ingredients in
Compare to active ingredients in	• •
Maximum Strength Mucinex®	-
SINUS-MAX® Day*	maximum strength
maximum strength	nighttime sinus
daytime sinus	acetaminophen
acetaminophen	diphenhydramine HCl,
guaifenesin, phenylephrine HCl	phenylephrine HCl
pain reliever, expectorant,	pain reliever,
nasal decongestant	antihistamine/cough suppressant,
ACTUAL	nasal decongestant
SIZE	ACTUAL
AGES	SIZE
12+	AGES
YEARS	12+
12	YEARS
CAPLETS	8
up&up™	CAPLETS
	up&up™
12 DAY AI	ND 8 NIGHT

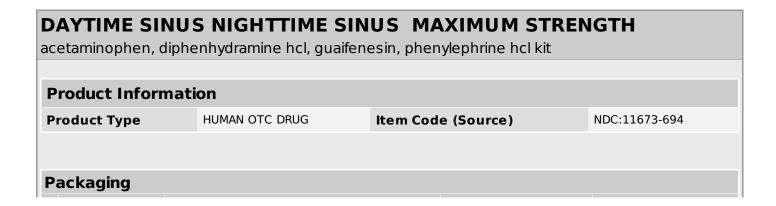
TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

*This product is not manufactured or distributed by Reckitt Benckiser LLC, owner of the registered trademark Maximum Strength Mucinex® SINUS-MAX® Day & Night. 50844 REV0517B61569409

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Target 44-615694



	ltem Code	Pack	age Description	M	arketing Star Date		ting End ate
		1 in 1 CARTON; Product	Type 0: Not a Combi	nation 07/0	1/2017		
	-						
Qu	antity of Pa	arts					
Par	t #	Package Qu	antity		Total Produc	t Quantity	
Part	1 BLISTER	РАСК		12			
Part	1 BLISTER	РАСК		8			
Pa	rt 1 of 2						
DA		INUS MAX		ENGTH			
ace	taminophen,	quaifenesin, r	henylephrine hcl	l tablet, film c	oated		
D							
Pro	oduct Infor	mation					
Rou	ite of Admini	stration C	ORAL				
Act	ive Inaredi	ent/Active M	oietv				
			ent Name		Basis o	of Strength	Strengt
ACE	TAMINOPHEN		D) (ACETAMINOPHEN	- UNII: 36209ITI 9		-	325 mg
		•	GUAIFENESIN - UNII:4		GUAIFENES		200 mg
			(UNII: 04JA59TNSJ) (- PHENYLEPH	RINE	
	1WS297W6MV)				HYDROCHLO	ORIDE	5 mg
	ctive Ingre	dianta					
Ina	clive mare	alents					
Ina			In a word is wet No.				
	-		Ingredient Na	me		S	trength
STA	RCH, CORN (UN	NII: 08232NY3SJ)		me		S	strength
STA CRO	RCH, CORN (UN SPOVIDONE, U	VII: 08232NY3SJ) JNSPECIFIED (UI	NII: 2S7830E561)	me		S	Strength
STA CRO FD&	RCH, CORN (UN SPOVIDONE, U C RED NO. 40	NII: 08232NY3SJ) JNSPECIFIED (UI (UNII: WZB9127X	NII: 2S7830E561) OA)	me		S	Strength
STA CRO FD& FD&	RCH, CORN (UP SPOVIDONE, U C RED NO. 40 C YELLOW NO	NII: 08232NY3SJ) JNSPECIFIED (UI (UNII: WZB9127X 9.6 (UNII: H77VEI	NII: 257830E561) OA) 93A8)	me			Strength
STA CRO FD& FD& MAG	RCH, CORN (UN SPOVIDONE, U C RED NO. 40 C YELLOW NO GNESIUM STEA	NII: 08232NY3SJ) JNSPECIFIED (UI (UNII: WZB9127X 9.6 (UNII: H77VEI RATE (UNII: 7009	NII: 257830E561) OA) 93A8) 7M6I30)	me		<u>ح</u>	Strength
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STA CRO FD& FD& MAG MAL MIC	RCH, CORN (UP SPOVIDONE, U C RED NO. 40 C YELLOW NO GNESIUM STEA TODEXTRIN (U ROCRYSTALLIN	NII: 08232NY3SJ) JNSPECIFIED (UI (UNII: WZ B9127X 0.6 (UNII: H77VEI RATE (UNII: 7009 NII: 7CVR7L4A2D) NE CELLULOSE (NII: 2S7830E561) OA) 93A8) 7M6I30) UNII: OP1R32D61U)			۲	Strength
STA CRO FD& MAG MAG MAL MIC POL	RCH, CORN (UN SPOVIDONE, U C RED NO. 40 C YELLOW NO GNESIUM STEA TODEXTRIN (U ROCRYSTALLIN YETHYLENE GI	NII: 08232NY3SJ) JNSPECIFIED (UI (UNII: WZ B9127X 0. 6 (UNII: H77VEI RATE (UNII: 7009 NII: 7CVR7L4A2D) NE CELLULOSE (LYCOL, UNSPEC	NII: 257830E561) OA) 93A8) 7M6I30) UNII: OP1R32D61U) IFIED (UNII: 3WJQ051	DW1A)			Strength
STA CRO FD& FD& MAG MAL MIC POL	RCH, CORN (UP SPOVIDONE, U C RED NO. 40 C YELLOW NO GNESIUM STEA TODEXTRIN (U ROCRYSTALLIN YETHYLENE GI	NII: 08232NY3SJ) JNSPECIFIED (UI (UNII: WZ B9127X 0.6 (UNII: H77VEI RATE (UNII: 7009 NII: 7CVR7L4A2D) NE CELLULOSE (LYCOL, UNSPECIFII	NII: 257830E561) OA) 93A8) 7M6I30) UNII: OP1R32D61U) IFIED (UNII: 3WJQ051 ED (UNII: 532B59J990	DW1A)		۲	Strength
STA CRO FD& FD& MAG MAL MIC POL POL	RCH, CORN (UN SPOVIDONE, U C RED NO. 40 C YELLOW NO GNESIUM STEA TODEXTRIN (U ROCRYSTALLIN YETHYLENE GI YVINYL ALCOH	NII: O8232NY3SJ) JNSPECIFIED (UI (UNII: WZ B9127X O. 6 (UNII: H77VE!! RATE (UNII: 7009 NII: 7CVR7L4A2D) NE CELLULOSE (LYCOL, UNSPECIFII CIFIED (UNII: FZ	NII: 2S7830E561) OA) 93A8) 7M6I30) UNII: OP1R32D61U) IFIED (UNII: 3WJQ0SI ED (UNII: 532B59J990 989GH94E)	DW1A)			Strength
STA CRO FD& MAC MAL MIC POL POL SILI	RCH, CORN (UP SPOVIDONE, U C RED NO. 40 C YELLOW NO GNESIUM STEA TODEXTRIN (U ROCRYSTALLIN YETHYLENE GI YVINYL ALCOH IDONE, UNSPE CON DIOXIDE (NII: O8232NY3SJ) JNSPECIFIED (UI (UNII: WZ B9127X 0.6 (UNII: H77VEI RATE (UNII: 7009 NII: 7CVR7L4A2D) NE CELLULOSE (LYCOL, UNSPECI IOL, UNSPECIFII CIFIED (UNII: FZ (UNII: ETJ7Z6XBU4	NII: 2S7830E561) OA) 93A8) 7M6I30) UNII: OP1R32D61U) IFIED (UNII: 3WJQ0SI ED (UNII: 532B59J990 989GH94E) 4)	DW1A) D)			Strength
STA CRO FD& FD& MAG MAL MIC POL POL SILIO SOD	RCH, CORN (UN SPOVIDONE, U C RED NO. 40 C YELLOW NO SNESIUM STEA TODEXTRIN (U ROCRYSTALLIN YETHYLENE GI YVINYL ALCOH IDONE, UNSPE CON DIOXIDE (DIUM STARCH C	NII: 08232NY3SJ) JNSPECIFIED (UI (UNII: WZ B9127X 0. 6 (UNII: H77VEI RATE (UNII: 7009 NII: 7CVR7L4A2D) NE CELLULOSE (LYCOL, UNSPECIFII CLYCOL, UNSPECIFII CIFIED (UNII: FZ CUNII: ETJ7Z 6XBU4 GLYCOLATE TYP	NII: 2S7830E561) OA) 93A8) 7M6I30) UNII: OP1R32D61U) IFIED (UNII: 3WJQ0SI ED (UNII: 532B59J990 989GH94E)	DW1A) D)			Strength
STA CRO FD& FD& MAL MIC POL POL SILIO SILIO SOD	RCH, CORN (UP SPOVIDONE, U C RED NO. 40 C YELLOW NO GNESIUM STEA TODEXTRIN (U ROCRYSTALLIN YETHYLENE GI YVINYL ALCOH IDONE, UNSPE CON DIOXIDE (NII: 08232NY3SJ) JNSPECIFIED (UI (UNII: WZ B9127X 0. 6 (UNII: H77VEI RATE (UNII: 7009 NII: 7CVR7L4A2D) NE CELLULOSE (LYCOL, UNSPECI IOL, UNSPECIFII CIFIED (UNII: FZ (UNII: ETJ7Z6XBU4 GLYCOLATE TYP I: 4ELV7Z65AP)	NII: 2S7830E561) OA) 93A8) 7M6I30) UNII: OP1R32D61U) IFIED (UNII: 3WJQ0SI ED (UNII: 532B59J990 989GH94E) 4)	DW1A) D)			Strength

Shape OVAL Size	no score 19mm 44;615 Marketing End
Shape OVAL Size Imprint Code Flavor Imprint Code 4 Contains Imprint Code 4 Packaging Imprint Code 4 Item Package Description Marketing Start Date 1 12 in 1 BLISTER PACK; Type 0: Not a Combination 14	19mm 44;615
Flavor Imprint Code 4 Contains 4 Package Description Marketing Start Date 1 12 in 1 BLISTER PACK; Type 0: Not a Combination	44;615
Contains Antipage Packaging Marketing Start # Item Code Package Description Marketing Start Date 1 12 in 1 BLISTER PACK; Type 0: Not a Combination Marketing Start	
Package Description Marketing Start Date Item Code Package Description 1 12 in 1 BLISTER PACK; Type 0: Not a Combination	Marketing End
# Item Code Package Description Marketing Start Date 1 12 in 1 BLISTER PACK; Type 0: Not a Combination	Marketing End
# Item Code Package Description Marketing Start Date 1 12 in 1 BLISTER PACK; Type 0: Not a Combination	Marketing End
# Item Code Package Description Marketing Start Date 1 12 in 1 BLISTER PACK; Type 0: Not a Combination	Marketing End
** Code Package Description Date 1 12 in 1 BLISTER PACK; Type 0: Not a Combination	Marketing End
	Date
Marketing Information	
Marketing CategoryApplication Number or Monograph CitationMarketing Start Date	Marketing End Date
OTC Monograph Drug M012 06/30/2013	
Part 2 of 2	
NIGHTTIME SINUS MAXIMUM STRENGTH	
acetaminophen, diphenhydramine hcl, phenylephrine hcl tablet, film coated	
Product Information	
Route of Administration ORAL	
Route of Administration ORAL	
Route of Administration ORAL Active Ingredient/Active Moiety	rongth Strongt
Route of Administration ORAL Active Ingredient/Active Moiety Ingredient Name Basis of Str	
Route of Administration ORAL Active Ingredient/Active Moiety Ingredient Name Basis of Str ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	325 mg
Route of Administration ORAL Active Ingredient/Active Moiety Ingredient Name Basis of Str ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) DIPHENHYDRAMINE	NE 12.5 mg
Route of Administration ORAL Active Ingredient/Active Moiety Basis of State Ingredient Name Basis of State ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) DIPHENHYDRAMINE - UNII:8GTS82S83M) PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - PHENYLEPHRINE -	325 mg
Route of Administration ORAL Active Ingredient/Active Moiety Basis of State Ingredient Name Basis of State ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) DIPHENHYDRAMINE - UNII:8GTS82S83M) PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - PHENYLEPHRINE -	325 mg
Route of Administration ORAL Active Ingredient/Active Moiety Basis of Stress of S	325 mg
Route of Administration ORAL Active Ingredient/Active Moiety Ingredient Name Basis of Str ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) DIPHENHYDRAMINE - UNII:8GTS82S83M) PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - PHENYLEPHRINE HYDROCHLORIDE INII: 04JA59TNSJ)	325 mg NE 12.5 mg
Route of Administration ORAL Active Ingredient/Active Moiety Active Ingredient/Active Moiety Basis of Str Active Ingredient Name Basis of Str Acetaminophen (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) DIPHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS 297 W6MV) Ingredient Name	325 mg
Route of Administration ORAL Active Ingredient/Active Moiety Active Ingredient/Active Moiety Ingredient Name Basis of Str ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) DIPHENHYDRAMINE - UNII:8GTS82S83M) PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - HYDROCHLORIDE PHENYLEPHRINE - PHENYLEPHRINE - HYDROCHLORIDE Inactive Ingredients Ingredient Name STARCH, CORN (UNII: 08232NY3SJ)	325 mg NE 12.5 mg
Route of Administration ORAL Active Ingredient/Active Moiety Basis of Stress of S	325 mg NE 12.5 mg
Route of Administration ORAL Active Ingredient/Active Moiety Basis of Str Acetaminophen (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) DIPHENHYDRAMINE - UNII:8GT582583M) PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:8297W6MV) PHENYLEPHRINE - UNII:04JA59TNSJ) (PHENYLEPHRINE - UNII:80CHLORIDE Inactive Ingredients Ingredient Name STARCH, CORN (UNII: 08232NY3SJ) CROSCARMELLOSE SODIUM (UNII: M280L1HH48) CROSPOVIDONE, UNSPECIFIED (UNII: 257830E561) Crossante Content Cont	325 mg NE 12.5 mg
Route of Administration ORAL Active Ingredient/Active Moiety Basis of Str Active Ingredient/Active Moiety Acteraminophen Name Basis of Str ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) DIPHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - Ingredients Ingredient Name STARCH, CORN (UNII: 08232NY3SJ) CROSCARMELLOSE SODIUM (UNII: M280L1HH48) CROSPOVIDONE, UNSPECIFIED (UNII: 257830E561) FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: 39EQA3S2JM)	325 mg NE 12.5 mg
Route of Administration ORAL Active Ingredient/Active Moiety Basis of Sti Active Ingredient/Active Moiety Active Ingredient Name Active Ingredient/Active Moiety Active Ingredient/Active Moiety Active Ingredient/Active Moiety Active Ingredient/Active Moiety Active Ingredients/Active Moiety DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) DIPHENHYDRAMINE - UNII:86T582583M) PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - Ingredients STARCH, CORN (UNII: 08232NY3SJ) CROSCARMELLOSE SODIUM (UNII: M280L1HH48) CROSPOVIDONE, UNSPECIFIED (UNII: 257830E561) FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: 3PEQA3S2JM) FD&C BLUE NO. 2 ALUMINUM LAKE (UNII: 4AQJ3LG584)	325 mg NE 12.5 mg
Route of Administration ORAL Active Ingredient/Active Moiety Basis of Strats AcetaMinophen (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) DIPHENHYDRAMINE - UNII:36TS82S83M) PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNICOCHLORIDE PHENYLEPHRINE - VINI:36TS82S83M) Inactive Ingredients Ingredient Name STARCH, CORN (UNII: 08232NY3SI) Ingredient Name STARCH, CORN (UNII: 08232NY3SI) CROSCARMELLOSE SODIUM (UNII: M280L1HH48) CROSPOVIDONE, UNSPECIFIED (UNII: 257830E561) FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	325 mg NE 12.5 mg

METHACRYLIC ACID AND ETHYL ACRYLATE COPOLYMER (UNII: NX76LV5T8J)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	blue	Score	no score
Shape	OVAL	Size	19mm
Flavor		Imprint Code	44;694
Contains			

Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC Monograph Drug	M012	07/01/2017	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC Monograph Drug	M012	07/01/2017	

Labeler - Target Corporation (006961700)

Establishment				
Name	Add	Iress	ID/FEI	Business Operations
LNK International, Inc.		038	154464	pack(11673-694)
Establishment				
Nama	Addrocc			Business Operations

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(11673-694) , pack(11673-694)

stablishment			
Name	Address	ID/FEI	Business Operations
K International, Inc.		832867894	manufacture(11673-694)
stablishment			
Name	Address	ID/FEI	Business Operations
K International, Inc.		967626305	pack(11673-694)
stablishment			
Name	Address	ID/FEI	Business Operations
		117025878	manufacture(11673-694)
Name	Address	-	

Revised: 8/2023

Target Corporation