TARGET MULTI SYMPTOM FLU RELIEF- acetaminophen, dextromethorphan hbr, and acetaminophen, chlorpheniramine maleate, dextromethorphan hbr TARGET CORPORATION

Up & up Multi-Symptom Flu Relief Combo Pack Drug Facts

Flu Relief Max Strength** Daytime Powder

Active ingredients (in each packet)

Acetaminophen 1000 mg

Dextromethorphan HBr 30 mg

Purposes

Pain reliever/Fever reducer

Cough suppressant

Uses

- temporarily relieves these symptoms due to a common cold or flu:
 - headache
 - minor aches and pains
 - cough due to minor throat and bronchial irritation
 - minor sore throat pain
- 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.

Sore throat warning: If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting consult a doctor promptly.

Do not use

- in a child under 12 years of age
- if you are allergic to acetaminophen
- 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

- liver disease
- a breathing problem such as emphysema or chronic bronchitis
- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, or emphysema

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

- pain 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 overdose, get medical help or contact a Poison Control Center right away at 1-800-222-1222. Prompt 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

• take every 6 hours, while symptoms persist. Do not take more than 3 packets in 24 hours unless directed by a doctor.

1. Age	1. Dose
1. adults and children 12 years of age and over	1. one packet

1.	children under 12 years of age	1. do not use

- dissolve contents of one packet into 8 oz. hot water; sip while hot. Consume the entire drink within 10-15 minutes.
- if using a microwave, add contents of one packet to 8 oz. of cool water; stir briskly before and after heating. Do not overheat.

Other information

1

- each packet contains: potassium 6 mg
- store at room temperature. Protect from excessive heat and moisture.

Inactive ingredients

anhydrous citric acid, caramel, flavor, maltodextrin, potassium chloride, silica, sucralose, sucrose.

Questions or comments?

1-866-467-2748

Flu Relief Max Strength** Nighttime Powder

Drug Facts

Active ingredients (in each packet)

Acetaminophen 1000 mg Chlorpheniramine maleate 4 mg Dextromethorphan HBr 30 mg

Purposes

Pain reliever/Fever reducer

Antihistamine

Cough suppressant

Uses

- temporarily relieves these symptoms due to a common cold or flu:
 - headache
 - minor aches and pains

- cough due to minor throat and bronchial irritation
- minor sore throat pain
- runny nose
- 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.

Sore throat warning: If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting consult a doctor promptly.

Do not use

- in a child under 12 years of age
- if you are allergic to acetaminophen
- 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 a MAO, ask a doctor or pharmacist before taking this product.

Ask a doctor before use if you have

- liver disease
- glaucoma
- trouble urinating due to an enlarged prostate gland
- a breathing problem such as emphysema or chronic bronchitis
- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, or emphysema

Ask a doctor or pharmacist before use if you are

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers

When using this product

avoid alcoholic drinks

- marked drowsiness may occur
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

Stop use and ask a doctor if

- pain 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 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.

Directions

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- if using a microwave, add contents of one packet to 8 oz. of cool water; stir briskly before and after heating. Do not overheat

Other information

- each packet contains: potassium 6 mg
- store at room temperature. Protect from excessive heat and moisture.

Inactive ingredients

anhydrous citric acid, caramel, flavor, maltodextrin, potassium chloride, silica, sucralose, sucrose.

Questions or comments?

1-866-467-2748

Other Safety Information

READ ALL WARNINGS AND DIRECTIONS ON CARTON BEFORE USE. KEEP CARTON FOR REFERENCE. DO NOT DISCARD.

DO NOT TAKE BOTH PRODUCTS AT THE SAME TIME.

DO NOT TAKE MORE THAN 3 DOSES IN TOTAL IN ANY 24 HOUR PERIOD.

DO NOT TAKE A DOSE OF THE NIGHTTIME PRODUCT SOONER THAN 4 HOURS AFTER THE LAST DOSE OF THE DAYTIME PRODUCT UNLESS DIRECTED BY YOUR DOCTOR.

TAMPER-EVIDENT INNER UNIT. DO NOT USE IF NECKBAND PRINTED WITH "SEALED FOR SAFETY" IS TORN OR MISSING.

Distributed by:

Principal Display Panel

Compare to the active ingredients in Theraflu Multi-Symptom Flu Relief Max Strength** Daytime* & Nighttime*.

NDC 82442-559-12

MULTI SYMPTOM FLU RELIEF

Daytime Maximum Strength** Flu Relief

AcetaminophenPain Reliever/Fever Reducer

Dextromethorphan HBrCough Suppressant

6 PACKETS

Nighttime Maximum Strength** Flu Relief

AcetaminophenPain Reliever/Fever Reducer

Chlorpheniramine MaleateAntihistamine

Dextromethorphan HBrCough Suppressant

6 PACKETS

Hot liquid therapy that relieves:

- Fever
- Body ache
- Headache
- Sore throat pain

- Cough
- Runny nose (Nighttime only)

6 DAYTIME PACKETS

6 NIGHTTIME PACKETS

12 TOTAL PACKETS

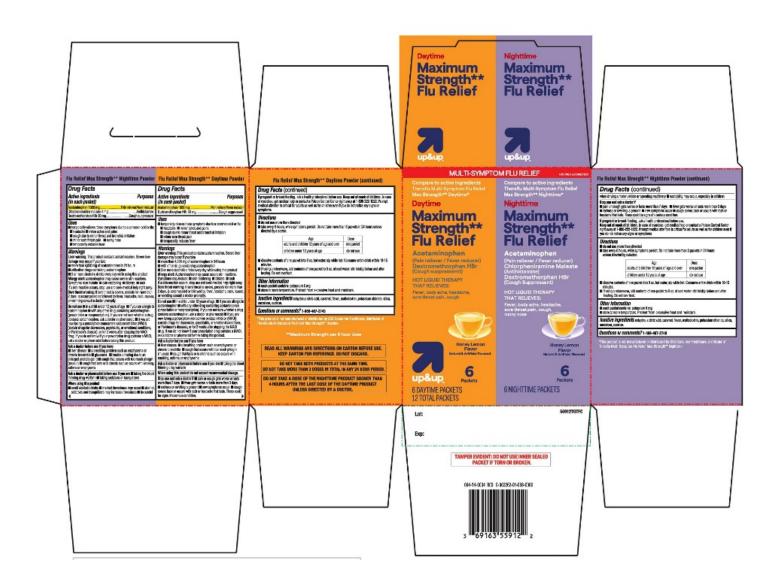
Honey Lemon Flavor

Natural & Artificial Flavored

*This product is not distributed by GSK Consumer Healthcare, distributor of Theraflu Multi-Symptom Flu Relief Max Strength** Daytime.

* This product is not distributed by GSK Consumer Healthcare, distributor of Theraflu Multi-Symptom Flu Relief Max Strength** Nighttime.

**Maximum Strength per 6 hour dose



TARGET MULTI SYMPTOM FLU RELIEF

acetaminophen, dextromethorphan hbr, and acetaminophen, chlorpheniramine maleate, dextromethorphan hbr kit

Product Infor	mation						
Product Type	HUMAN	OTC DRUG	Item Code	(Sourc	e) i	NDC:82442-	559
Packaging							
				Mark	eting Start	Market	ing End
# Item Code	Pa	ckage Descriptio	n	PIGIK	Date		ate
1 NDC:82442-559- 12	1 in 1 CARTON Package	; Type 1: Convenience	e Kit of Co-	06/06/20)25		
Quantity of Pa	arts						
Part #	Package (Quantity		Tota	al Product Qu	antity	
Part 1 1 CARTON			6				
Part 2 1 CARTON			6				
D							
Part 1 of 2							
TARGET D	AYTIME	MS FLU					
acetaminophen	dextrometh	orphan hbr powde	er				
•		· ·					
Product Infor	mation						
Item Code (Sou	rce)	NDC:82442-558					
Route of Admini	stration	ORAL					
Active Ingredi	ent/Active	Moiety					
	Ingree	dient Name			Basis of St	renath	
ACETAMINOPHEN	(UNII: 36209ITL					· • · · y · · ·	Strength
DEXTROMETHORP		9D) (ACETAMINOPHEN	I - UNII:36209	PITL9D)	ACETAMINOPHEN	-	Strength 1000 mg
(DEXTROMETHORPH		ROMIDE (UNII: 9D2RT			ACETAMINOPHEN DEXTROMETHORF HYDROBROMIDE	_	-
		ROMIDE (UNII: 9D2RT			DEXTROMETHOR	_	1000 mg
	AN - UNII:7355X	ROMIDE (UNII: 9D2RT			DEXTROMETHOR	_	1000 mg
(DEXTROMETHORPH	AN - UNII:7355X	ROMIDE (UNII: 9D2RT	'I9KYH)		DEXTROMETHOR	PHAN	1000 mg
(DEXTROMETHORPH	AN - UNII:7355X	ROMIDE (UNII: 9D2RT (3ROTS) Ingredient Nam	'I9KYH)		DEXTROMETHOR	PHAN	1000 mg 30 mg
(DEXTROMETHORPH	AN - UNII:7355X dients IC ACID (UNII: 3	ROMIDE (UNII: 9D2RT (3ROTS) Ingredient Nam	'I9KYH)		DEXTROMETHOR	PHAN	1000 mg 30 mg
(DEXTROMETHORPH	AN - UNII:7355X : dients IC ACID (UNII: 2 D99G2B1R)	ROMIDE (UNII: 9D2RT (3ROTS) Ingredient Nam (XF417D3PSL)	'I9KYH)		DEXTROMETHOR	PHAN	1000 mg 30 mg
(DEXTROMETHORPH Inactive Ingre ANHYDROUS CITR CARAMEL (UNII: T9	AN - UNII:7355X Adients IC ACID (UNII: 2 D99G2B1R) JNII: 7CVR7L4A2	ROMIDE (UNII: 9D2RT (3ROTS) Ingredient Nam XF417D3PSL) D)	'I9KYH)		DEXTROMETHOR	PHAN	1000 mg 30 mg
(DEXTROMETHORPH Inactive Ingre ANHYDROUS CITR CARAMEL (UNII: T9 MALTODEXTRIN (U	AN - UNII:7355X idients IC ACID (UNII: 2 D99G2B1R) JNII: 7CVR7L4A2 RIDE (UNII: 660 (UNII: ETJ7Z6XB	ROMIDE (UNII: 9D2RT (3ROTS) Ingredient Nam XF417D3PSL) D))YQ98I10)	'I9KYH)		DEXTROMETHOR	PHAN	30 mg

Product Cha	ractoristics					
		te, yellow, beige, and brown color)		Scol	re	
Shape		, ,,		Size	-	
-	HONEY (Lemon)			Impi	rint Code	•
Contains						
Packaging						
# Item Code	e Pa	ackage Description	Mar	keting Start Date		ting End ate
1 NDC:82442-55	8- 6 in 1 CARTO Package	N; Type 1: Convenience Kit of Co-				
Marketing	g Informa	tion				
Marketing Category	Applic	ation Number or Monograph Citation	Ma	rketing Start Date		eting End Date
OTC Monograph I	Drug M012		06/06	5/2025		
	2					
	NIGHTTIN	IE MS FLU RELIEF ramine maleate, dextromethor	phan l	hbr powder		
TARGET	NIGHTTIN		phan	hbr powder		
TARGET acetaminophe	NIGHTTIN en, chlorpheni		phan l	hbr powder		
TARGET acetaminophe Product Infe	NIGHTTIN en, chlorpheni ormation		phan l	hbr powder		
TARGET	NIGHTTIN en, chlorpheni ormation ource)	ramine maleate, dextromethor	phan l	hbr powder		
TARGET acetaminophe Product Infe Item Code (So	NIGHTTIN en, chlorpheni ormation ource)	ramine maleate, dextromethor NDC:82442-560	phan l	hbr powder		
TARGET acetaminophe Product Infe Item Code (So Route of Adm	NIGHTTIN en, chlorpheni ormation ource) inistration	ramine maleate, dextromethor NDC:82442-560 ORAL	phan I	hbr powder		
TARGET acetaminophe Product Infe Item Code (So Route of Adm Active Ingre	NIGHTTIN en, chlorpheni ormation ource) inistration dient/Active	ramine maleate, dextromethor NDC:82442-560 ORAL MOiety edient Name		hbr powder Basis of St	rength	Strengt
TARGET acetaminophe Product Info Item Code (So Route of Adm Active Ingre	NIGHTTIN en, chlorpheni ormation ource) inistration dient/Active Ingre	ramine maleate, dextromethor NDC:82442-560 ORAL MOiety dient Name FL9D) (ACETAMINOPHEN - UNII:36209)	TL9D)	Basis of Stu ACETAMINOPHEN	_	Strengt 1000 mg
TARGET acetaminophe Product Info Item Code (So Route of Adm Active Ingre ACETAMINOPHE CHLORPHENIRA	NIGHTTIN en, chlorpheni ormation ource) inistration dient/Active Ingre SN (UNII: 362091 MINE MALEATE	ramine maleate, dextromethor NDC:82442-560 ORAL MOiety edient Name	TL9D)	Basis of St	_	-
TARGET acetaminophe Product Info Item Code (So Route of Adm Active Ingre ACETAMINOPHE CHLORPHENIRA UNII:3U6I01965U DEXTROMETHO	NIGHTTIN en, chlorpheni ormation ource) inistration dient/Active Ingre EN (UNII: 362091 MINE MALEATE	ramine maleate, dextromethor NDC:82442-560 ORAL Moiety edient Name FL9D) (ACETAMINOPHEN - UNII:36209I (UNII: V1Q0090J9Z) (CHLORPHENIRA BROMIDE (UNII: 9D2RTI9KYH)	TL9D)	Basis of Stu ACETAMINOPHEN CHLORPHENIRAMI	NE	1000 mg
TARGET acetaminophe Product Infe Item Code (So Route of Adm Active Ingre ACETAMINOPHE CHLORPHENIRA UNII:3U6I01965U DEXTROMETHOR (DEXTROMETHOR	NIGHTTIN en, chlorpheni ormation ource) inistration dient/Active ingra N(UNII: 362091 MINE MALEATE N RPHAN HYDROI PHAN - UNII:7355	ramine maleate, dextromethor NDC:82442-560 ORAL Moiety edient Name FL9D) (ACETAMINOPHEN - UNII:36209I (UNII: V1Q0090J9Z) (CHLORPHENIRA BROMIDE (UNII: 9D2RTI9KYH)	TL9D)	Basis of Stu ACETAMINOPHEN CHLORPHENIRAMI MALEATE DEXTROMETHORP	NE	1000 mg 4 mg
TARGET acetaminophe Product Infe Item Code (So Route of Adm Active Ingre ACETAMINOPHE CHLORPHENIRA UNII:3U6I01965U DEXTROMETHOR (DEXTROMETHOR	NIGHTTIN en, chlorpheni ormation ource) inistration dient/Active ingra N(UNII: 362091 MINE MALEATE N RPHAN HYDROI PHAN - UNII:7355	ramine maleate, dextromethor NDC:82442-560 ORAL MOiety edient Name FL9D) (ACETAMINOPHEN - UNII:362091 (UNII: V1Q0090J9Z) (CHLORPHENIRA BROMIDE (UNII: 9D2RTI9KYH) SX3ROTS)	TL9D)	Basis of Stu ACETAMINOPHEN CHLORPHENIRAMI MALEATE DEXTROMETHORP	NE PHAN	1000 mg 4 mg 30 mg
TARGET acetaminophe Product Infe Item Code (So Route of Adm Active Ingre ACETAMINOPHE CHLORPHENIRA UNII: 3U6I01965U	NIGHTTIN en, chlorpheni ormation ource) inistration dient/Active Ingre EN (UNII: 362091 MINE MALEATE MINE MALEATE PHAN - UNII:7355	ramine maleate, dextromethor NDC:82442-560 ORAL e Moiety edient Name FL9D) (ACETAMINOPHEN - UNII:36209I (UNII: V1Q0090J9Z) (CHLORPHENIRA BROMIDE (UNII: 9D2RTI9KYH) SX3ROTS) Ingredient Name	TL9D)	Basis of Stu ACETAMINOPHEN CHLORPHENIRAMI MALEATE DEXTROMETHORP	NE PHAN	1000 mg 4 mg

	CHIOR	IDE (UNII: 660YQ98I10)			
		NII: ETJ7Z 6XBU4)			
SUCRALOSE					
SUCROSE (U					
Product (Charac	teristics			
Color white (white to off-white, yellow, beige, and brown color)		olor)	Score		
Shape				Size	
Flavor	HONE	Y (lemon)		Imprint Code	
Contains					
Packagin	g				
# Item C	ode	Package Description	Marketing Start Date	Marketing E Date	nd
				Date	
1 NDC:82442		5 in 1 CARTON; Type 0: Not a Combination Product	2000		
• 06	I	Product			
• 06	ing Ir		Marketing Sta Date		nd
Market Market Catego	ing II	Product Iformation Application Number or Monograph Citation	.	rt Marketing E	nd
Market Market Catego	ing II	Product Iformation Application Number or Monograph Citation	Date	rt Marketing E	nd
Market Market Catego OTC Monogra	ing II ting ory aph Drug	formation Application Number or Monograph Citation	Date	rt Marketing E	nd
Market Market Catego OTC Monogra	ing II ting ory aph Drug	Product Iformation Application Number or Monograph Citation	Date	rt Marketing E	nd
Market Market Catego OTC Monogra	ing II ting ory aph Drug ing II	formation Application Number or Monograph Citation	Date	rt Marketing E Date	

Labeler - TARGET CORPORATION (006961700)

Revised: 6/2025

TARGET CORPORATION