THERAFLU FLU RELIEF MAX STRENGTH NIGHTTIME- acetaminophen, chlorpheniramine maleate, dextromethorphan hbr powder Haleon US Holdings LLC

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 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
- glaucoma
- trouble urinating due to an enlarged prostate gland
- 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

- 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.

Age	Dose
adults and children	one packet
12 years of age and over	
children under	do not use
12 years of age	

- dissolve contents of one packet into 8 oz. hot water; sip while hot. Consume 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

- each packet contains:potassium 5 mg, sodium 22 mg
- phenylketonurics:contains phenylalanine 12.9 mg per packet
- store at a controlled room temperature at 20 25°C (68 77°F)

Inactive ingredients

acesulfame potassium, anhydrous citric acid, aspartame, D&C yellow no. 10, FD&C blue no. 1, FD&C red no. 40, maltodextrin, natural and artificial flavors, silicon dioxide, sodium citrate, soy lecithin, sucrose, tribasic calcium phosphate

Questions or comments?

1-855-297-3031

Additional Information

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

HOT LIQUID THERAPYthat relieves:

Fever

Body Ache

Headache

Sore Throat Pain

Cough

Runny Nose

Honey Lemon Flavor

*Maximum Strength per 6 hour dose

**Temporarily controls cough to help you rest. This is not a sleep-aid.

PARENTS:Learn about teen medicine abuse

www.StopMedicineAbuse.org

TAMPER EVIDENT INNER UNIT DO NOT USE IF SEALED THERAFLU PACKET IS TORN OR BROKEN

1-855-297-3031

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Principal Display Panel

MULTI-SYMPTOM FLU RELIEF

HALEON

THERAFLU

FLU RELIEF MAX STRENGTH*

NIGHTTIME

Acetaminophen

Pain Reliever/Fever Reducer

Chlorpheniramine Maleate

Antihistamine

Dextromethorphan HBr

Cough Suppressant

HELPS YOU REST**

Hot liquid therapy that relieves:

/ Fever

/ Body ache

/ Headache

/ Sore throat pain

/ Runny nose

Honey Lemon Flavor

6 PACKETS



THERAFLU FLU RELIEF MAX STRENGTH NIGHTTIME

acetaminophen, chlorpheniramine maleate, dextromethorphan hbr powder

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0067-7922
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	1000 mg	
CHLORPHENIRAMINE MALEATE (UNII: V1Q0090J9Z) (CHLORPHENIRAMINE - UNII: 3U6I01965U)	CHLORPHENIRAMINE MALEATE	4 mg	
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg	

Inactive Ingredients		
Ingredient Name	Strength	
ACESULFAME POTASSIUM (UNII: 230V73Q5G9)		
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)		
ASPARTAME (UNII: Z0H242BBR1)		
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		
MALTODEXTRIN (UNII: 7CVR7L4A2D)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)		
SOYBEAN LECITHIN (UNII: 1DI56QDM62)		
SUCROSE (UNII: C151H8M554)		
TRIBASIC CALCIUM PHOSPHATE (UNII: 91D9GV0Z28)		

Product Characteristics			
Color	white (to off white, yellow, beige, and brown color)	Score	
Shape		Size	
Flavor	HONEY (Lemon)	Imprint Code	
Contains			

Packaging				
	# Item Co	le Package Description	Marketing Start Date	Marketing End Date
	1 NDC:0067-79	6 in 1 CARTON; Type 0: Not a Combinatio Product	n 06/15/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	06/15/2022	

Labeler - Haleon US Holdings LLC (079944263)

Revised: 5/2025 Haleon US Holdings LLC