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

Drug Facts

Active ingredients (in each 30 mL)

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:
    - 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
- 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

- **do not use more than directed**
- measure the dose correctly using the enclosed dosing cup
- adults and children 12 years of age and over: take every 6 hours in dosing cup provided, while symptoms persist
- do not take more than 3 doses (90 mL) in 24 hours unless directed by a doctor
- children under 12 years of age: do not use

Age	Dose
adults and children 12 years of age and over	30 mL
children under 12 years of age	do not use

Other information

- **each 30 mL contains**: potassium 5 mg, sodium 34 mg
- store at controlled room temperature 20-25°C (68-77°F)

Inactive ingredients

anhydrous citric acid, carboxymethylcellulose sodium, glycerin, nautral and artificial flavors, natural grade A honey, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate, sodium gluconate, sucralose, xanthan gum, zinc gluconate

Questions or comments?

1-855-328-5259

Additional Information

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

PEEL BACK HERE

DO NOT USE IF NECKBAND PRINTED WITH “SEALED FOR SAFETY” IS TORN OR MISSING.

***Maximum Strength per 6 hour dose.**

PARENTS: Learn about teen medicine abuse

www.StopMedicineAbuse.org

Principal Display Panel

THERAFLU

FLU RELIEF

MAX STRENGTH*

NIGHTTIME

Acetaminophen

Pain Reliever/Fever Reducer

Chlorpheniramine Maleate

Antihistamine

Dextromethorphan HBr

Cough Suppressant

HELPS YOU REST**

Powerful *fever fighting*formula that relieves:

/ Body ache

/ Headache

/ Sore throat pain

/ Cough

/ Runny nose

Honey & Elderberry Flavor

8.3 FL OZ (245.5 mL)

THERAFLU FLU RELIEF MAX STRENGTH NIGHTTIME

acetaminophen, chlorpheniramine maleate, dextromethorphan hbr syrup

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0067-8205
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	1000 mg

ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	in 30 mL
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	4 mg in 30 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
GLYCERIN (UNII: PDC6A3C0OX)	
HONEY (UNII: Y9H1V576FH)	
PEG-10 .BETA.-SITOSTERYL ETHER (UNII: B2138XJ83G)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SODIUM GLUCONATE (UNII: R6Q3791S76)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ZINC GLUCONATE (UNII: U6WSN5SQ1Z)	

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
May contain	**ANHYDROUS CITRIC ACID** (UNII: XF417D3PSL)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	BERRY (Elderberry)	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0067-8205-01	245.5 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package	06/27/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	06/27/2022	

Labeler - Haleon US Holdings LLC (079944263)