THERAFLU-D FLU RELIEF MAX STRENGTH PLUS NASAL DECONGESTANT NIGHTTIME- acetaminophen, chlorpheniramine maleate, dextromethorphan hbr, pseudoephedrine hcl syrup Haleon US Holdings LLC

Drug Facts

Active ingredients (in each 30 mL)

Acetaminophen 1000 mg

Chlorpheniramine maleate 4 mg

Dextromethorphan HBr 30 mg

Pseudoephedrine HCl 60 mg

Purposes

Pain reliever/Fever reducer

Antihistamine

Cough suppressant

Nasal decongestant

Uses

- temporarily relieves these symptoms due to a common cold or flu:
 - headache
 - minor aches and pains
 - minor sore throat pain
 - runny nose
 - cough due to minor throat and bronchial irritation
 - nasal and sinus congestion
- 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
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- 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

- do not exceed recommended dosage
- 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

- nervousness, dizziness, or sleeplessness occurs
- pain, cough 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
- 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 every 6 hours
•	children under 12 years of age	•	do not use

Other information

- each 30 mL contains: sodium 26 mg
- store at 20-25°C (68-77°F). Do not refrigerate. Keep bottle tightly closed.

Inactive ingredients

alcohol, anhydrous citric acid, carboxymethylcellulose sodium, D&C red no. 33, FD&C red no. 40, glycerin, natural and artificial flavors, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose

Questions or comments?

1-855-328-5259

Additional Information

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

PARENTS:

Learn about teen medicine abuse

www.StopMedicineAbuse.org

DO NOT USE IF TEAR BAND ON CAP IS BROKEN OR MISSING.

Distributed by: Haleon

Warren, NJ 07059

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Pat. Info www.productpats.com

Principal Display Panel

MULTI-SYMPTOM RELIEF NEWHALEON

THERAFLU-D

FLU RELIEF

MAX STRENGTH*

+ NASAL DECONGESTANT

NIGHTTIME

Acetaminophen

Pain Reliever/Fever Reducer

Chlorpheniramine Maleate

Antihistamine

Dextromethorphan HBr

Cough Suppressant

Pseudoephedrine HCI

Nasal Decongestant

Fast, Powerful Multi-Symptom relief of:

/ Fever

/ Nasal congestion

/ Sinus Pressure

/ Head & Body Ache

/ Sore Throat Pain

/ Cough

/ Runny Nose

/ Sneezing

Cherry Flavor

8.0 FL OZ (237 mL) 62000000209276 Carton Front



THERAFLU-D FLU RELIEF MAX STRENGTH PLUS NASAL DECONGESTANT NIGHTTIME

acetaminophen, chlorpheniramine maleate, dextromethorphan hbr, pseudoephedrine hcl syrup

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0067-0114
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	1000 mg in 30 mL		
CHLORPHENIRAMINE MALEATE (UNII: V1Q0090J9Z) (CHLORPHENIRAMINE - UNII:3U6I01965U)	CHLORPHENIRAMINE MALEATE	4 mg in 30 mL		
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg in 30 mL		
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	60 mg in 30 mL		

Inactive Ingredients				
Ingredient Name	Strength			
ALCOHOL (UNII: 3K9958V90M)				
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)				
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311)				
D&C RED NO. 33 (UNII: 9DBA0SBB0L)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
GLYCERIN (UNII: PDC6A3C0OX)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)				
SORBITOL (UNII: 506T60A25R)				
SUCRALOSE (UNII: 96K6UQ3ZD4)				

Product Characteristics			
Color	red (Clear)	Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0067- 0114-01	1 in 1 CARTON	06/19/2024	
1		237 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	06/19/2024	

Labeler - Haleon US Holdings LLC (079944263)

Revised: 5/2024 Haleon US Holdings LLC