

**THERAFLU SEVERE COLD RELIEF NIGHTTIME- acetaminophen,
diphenhydramine hcl syrup
Haleon US Holdings LLC**

Drug Facts

Active ingredients (in each 30 mL)

Acetaminophen 650 mg

Diphenhydramine HCl 25 mg

Purposes

Pain reliever/Fever reducer

Antihistamine/Cough suppressant

Uses

- temporarily relieves these symptoms to a cold:
 - minor aches and pains
 - minor sore throat pain
 - headache
 - runny nose
 - sneezing
 - itchy nose or throat
 - itchy, watery eyes due to hay fever
 - cough due to minor throat and bronchial irritation
- 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

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.
- with any other product containing diphenhydramine, even one used on the skin
- 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 sedatives or tranquilizers
- taking the blood thinning drug warfarin

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

- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- pain or cough gets worse or lasts more than 7 days
- 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
- take every 4 hours in dosing cup provided, while symptoms persist
- do not take more than 5 doses (150 mL) in 24 hours unless directed by a doctor

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 30 mg, sodium 13 mg
- store at controlled room temperature 20°-25°C (68° - 77°F)

Inactive ingredients

acesulfame potassium, anhydrous citric acid, edetate disodium, FD&C blue no. 1, FD&C red no. 40, glycerin, maltitol solution, natural and artificial flavors, propylene glycol, purified water, sodium benzoate, sodium citrate

Questions or comments?

1-855-328-5259

Other Information

Distributed by: Haleon, Warren, NJ 07059 Made in Canada Trademarks are owned by or licensed to the Haleon group of companies. ©2023 Haleon group of companies or its licensor. Pat. Info www.productpats.com

PEEL BACK HERE

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

62000000207729

Package/Label Principal Display Panel

MULTI-SYMPTOM **COLD RELIEF**

NEW FORMULA

THERAFLU

SEVERE

COLD RELIEF

NIGHTTIME

Acetaminophen

Pain Reliever/Fever Reducer

Diphenhydramine HCl

Antihistamine/Cough Suppressant

HELPS YOU REST*

Powerful formula that relieves:

/ Cough

/ Sore throat pain

/ Head and body ache

/ Fever

/ Runny nose

/ Sneezing

8.3 FL OZ (245.5 mL)

Berry Flavor

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

62000000207728

MULTI-SYMP TOM COLD RELIEF

NEW FORMULA

HALEON

THERAFLU

SEVERE COLD RELIEF

NIGHTTIME

Acetaminophen

Pain Reliever/Fever Reducer

Diphenhydramine HCl

Antihistamine/Cough Suppressant



Powerful formula that relieves:

- / Cough
- / Sore throat pain
- / Head and body ache
- / Fever
- / Runny nose
- / Sneezing

8.3 FL OZ (245.5 mL)

Berry Flavor

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

62000000207728



THERAFLU SEVERE COLD RELIEF NIGHTTIME

acetaminophen, diphenhydramine hcl syrup

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0067-0105
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg in 30 mL
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
ACESULFAME POTASSIUM (UNII: 230V73Q5G9)	

ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
MALTITOL (UNII: D65DG142WK)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	

Product Characteristics

Color	red	Score	
Shape		Size	
Flavor	BERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0067-0105-08	245.5 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/01/2023	

Labeler - Haleon US Holdings LLC (079944263)