FLU AND SEVERE COLD AND COUGH NIGHT TIME- acetaminophen, diphenhydramine hydrochloride, phenylephrine hydrochloride powder, for solution

Meijer Distribution Inc

Meijer Distribution, Inc. Flu & Severe Cold & Cough Drug Facts

Active ingredients (in each packet)

Acetaminophen 650 mg

Diphenhydramine hydrochloride 25 mg

Phenylephrine HCl 10 mg

Purposes

Pain reliever/fever reducer

Antihistamine/cough suppressant

Nasal decongestant

Uses

- temporarily relieves these symptoms due to a cold:
- minor aches and pains
- minor sore throat pain
- headache
- nasal and sinus congestion
- 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

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 have ever had an allergic reaction to this product or any of its ingredients
- 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 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
- 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 sedatives or tranquilizers
- taking the blood thinning drug warfarin

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
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- pain, cough or nasal congestion 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.

Overdose warning: In case of overdose, get medical help or contact a Poison Control Center right away (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 (see overdose warning)
- take every 4 hours, while symptoms persist. Do not take more than 5 packets in 24 hours unless directed by a doctor.

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

- 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 10 mg and sodium 25 mg
- **phenylketonurics:** contains phenylalanine 13 mg per packet
- store at 20-25°C (68-77°F). Protect product from heat and moisture.

Inactive ingredients

acesulfame potassium, anhydrous citric acid, aspartame, colloidal silicon dioxide, D&C yellow #10, FD&C blue #1, FD&C red #40, flavors, maltodextrin, pregelatinized starch, sodium citrate, sucrose, tribasic calcium phosphate

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

meijer_®

Compare to Theraflu® Nighttime Severe Cold & Cough active ingredients

NIGHTTIME flu & severe cold & cough

Acetaminophen | Pain Reliever | Fever Reducer

Diphenhydramine HCI | Antihistamine | Cough Suppressant

Phenylephrine HCl | Nasal Decongestant

Hot Liquid Therapy for Fast Relief

Multi-Symptom

Nasal & Sinus Congestion

Sore Throat Pain | Cough

Headache | Body Ache | Fever

Runny Nose | Sneezing

Alcohol-Free

Honey Lemon Infused with White Tea Flavors

6 Packets









DIST. BY MELJER DISTRIBUTION, INC. GRAND RAPIDS, MI 495 44 www.meijer.com o obxeM niebsM



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flu ક severe cold & couah

Hot Liquid Therapy for Fast Relief

Multi-Symptom

6 Packets

Drug Facts Active ingredients (in each packet)

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Warmings

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Drug Facts (continued)

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professional below reprofessional below reprofessional below reprofessional below rekape out of reach of children. Overdose
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Other information

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This product is not manufactured or distributed y Novartis Consumer Health S.A., owner of the egistered trademark Theraflu®.

FLU AND SEVERE COLD AND COUGH NIGHT TIME

acetaminophen, diphenhydramine hydrochloride, phenylephrine hydrochloride powder, for solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:41250-964

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) **ACETAMINOPHEN** 650 mg **DIPHENHYDRAMINE HYDROCHLORIDE** (UNII: TC2D6IAD40) DIPHENHYDRAMINE 25 mg (DIPHENHYDRAMINE - UNII:8GTS82S83M) **HYDROCHLORIDE**

PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE -PHENYLEPHRINE 10 mg UNII:1WS297W6MV) **HYDROCHLORIDE**

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

Product Characteristics				
Color	WHITE (mixture of white, light yellow-orange particles) , ORANGE	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:41250-964- 91	6 in 1 CARTON; Type 0: Not a Combination Product	09/03/2010	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	09/03/2010	

Labeler - Meijer Distribution Inc (006959555)

Revised: 10/2023 Meijer Distribution Inc