FLU AND SEVERE COLD AND COUGH NIGHTTIME- acetaminophen, diphenhydramine hydrochloride, phenylephrine hydrochloride powder, for solution Kroger Company

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Kroger Co. NightTime Flu plus Severe Cold & Cough Drug Facts

Active ingredients (in each packet)

Acetaminophen 650 mg Diphenhydramine HCl 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

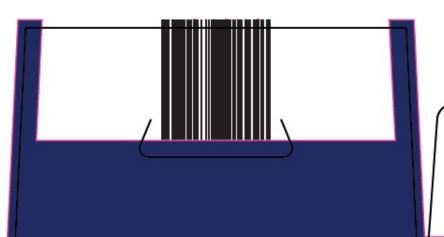
Questions or comments?

1-800-632-6900

Package/Label Principal Display Panel

COMPARE TO the active ingredients of THERAFLU[®] NIGHTTIME SEVERE COLD & COUGH

See side panel OUR PHARMACIST RECOMMENDED NightTime Flu + Severe Cold & Cough Acetaminophen Pain Reliever/Fever Reducer Diphenhydramine HCl Antihistamine/Cough Suppressant Phenylephrine HCl Nasal Decongestant Cough Nasal Congestion Sore Throat Pain Headache Body Ache Fever Runny Nose Sneezing Honey Lemon Flavor Infused with white tea flavors 6 PACKETS





COMPARE TO the active ingredients of THERAFLU® NIGHTTIME SEVERE COLD & COUGH *See side panel

NDC 30142-964-91



NightTime (** ^{**} Flu + Severe Cold & Cough

Acetaminophen Pain Reliever/Fever Reducer

Diphenhydramine HCl Antihistamine/ Cough Suppressant Phenylephrine HCl Nasal Decongestant

Cough Nasal Congestion Sore Throat Pain Headache Body Ache Fever Runny Nose Sneezing

Honey Lemon Flavor Infused with white tea flavors





NightTime (↓↓ Flu + Severe Cold & Cough

Acetaminophen Pain Reliever/Fever Reducer Diphenhydramine HCl Antihistamine/ Cough Suppressant Phenylephrine HCl Nasal Decongestant

> Cough Nasal Congestion Sore Throat Pain Headache Body Ache Fever Runny Nose Sneezing Honey Lemon Flavor Infused with white tea flavors



CONVENIENT RECLOSING TAB

DO NOT USE IF PRINTED PACKETS ARE TORN OR PUNCTURED
Drug Facts
Active ingredients (in each packet) Purposes
Acetaminophen 650 mgPain relieven/fever reducer
Diphenhydramine HCl 25 mg Phenylephrine HCl 10 mgNasal 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 fitchy, 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 vorniting, 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 Iver 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 glaucoma a breathing problem such as emphysema or chronic bronchitis cough that occurs with too much phlegm (mucus) glaucoma 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 taking the blood thinning drug warfarin taking the blood thinning drug warfarin
When using this product avoid alcoholic drinks 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 redness or swelling is present pain, cough or rasal congestion gets worse or lasts more than 3 days cough comes back or occurs with rash or headache that lasts. These could be signs of a serious condition.
OPEN OTHER END

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

Drug Facts (continued)

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Questions or comments? 1-800-632-6900

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FLU AND SEVERE COLD AND COUGH NIGHTTIME

acetaminophen, diphenhydramine hydrochloride, phenylephrine hydrochloride powder, for solution

Product In	formation					
Product Typ	e	HUMAN OTC DRUG	Item Code (S	ource)	NDC:30142	-964
Route of Adu	ministration	ORAL				
Active Ing	redient/Active Moi	etv				
		edient Name		Basis of St	trength	Strength
ACETAMINO	Ū.) (ACETAMINOPHEN - UNII:362	209ITL9D)	ACETAMINOPHEN	-	650 mg
	AMINE HYDROCHLOR	IDE (UNII: TC2D6JAD40) (DIPHI		DIPHENHYDRAMI HYDROCHLORIDE		25 mg
PHENYLEPHI UNII:1WS297V		E (UNII: 04JA59TNSJ) (PHENYLE	PHRINE -	PHENYLEPHRINE HYDROCHLORIDE	3	10 mg
Inactive In	gredients					
		Ingredient Name			S	trength
ACESULFAM	E POTASSIUM (UNII: 230	OV73Q5G9)				
ANHYDRO US	CITRIC ACID (UNII: XF4	17D3PSL)				
ASPARTAME	(UNII: Z0H242BBR1)					
SILICON DIO	XIDE (UNII: ETJ7Z6XBU	4)				
D&C YELLO	W NO. 10 (UNII: 35SW5U	SQ3G)				
FD&C BLUE	NO. 1 (UNII: H3R47K3TBI))				
FD&C RED N	O.40 (UNII: WZB9127XO	A)				
MALTODEXT	F RIN (UNII: 7CVR7L4A2D)				
SODIUM CIT	RATE, UNSPECIFIED FO	RM (UNII: 1Q73Q2JULR)				
SUCROSE (U	NII: C151H8M554)					
TRIBASIC CA	LCIUM PHO SPHATE (U	NII: 91D9GV0Z28)				
	haracteristics					
Color	WHITE (mixture of white	, light yellow-orange particles) ,	ORANGE		Score	
Shape					Size	
Flavor					Imprint Co	de
Contains						

# Item Code	Package Description	Marketing Start Date	Marketing End Date
NDC:30142-964-91	6 in 1 CARTON; Type 0: Not a Combination Product	09/02/2010	
Marketing Inf	e motion		
Marketing Info	ormation		
Marketing Info		Marketing Start Date	Marketing End Date

Labeler - Kroger Company (006999528)

Revised: 4/2020

Kroger Company