DAYTIME NIGHTTIME SEVERE COLD COUGH AND FLU RELIEF- acetaminophen, dextromethorphan hbr, phenylephrine hcl, diphenhydramine hcl Rite Aid Corporation

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.

Rite Aid Corporation Daytime Nighttime Severe Cold Cough & Flu Relief Drug Facts

Active ingredients (in each packet) - Daytime

Acetaminophen 500 mg

Dextromethorphan HBr 20 mg

Phenylephrine HCl 10 mg

Purpose

Pain reliever/fever reducer 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
- 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.
- 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
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, or emphysema
- a sodium-restricted diet

Ask a doctor or pharmacist before use if you are

taking the blood thinning drug warfarin

When using this product

do not exceed recommended dosage

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 6 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 22 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

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

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- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

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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.

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Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

COMBO PACK

Compare to the active ingredients of Theraflu® Severe Cold and Theraflu® Nighttime Severe Cold & Cough

daytime

severe cold cough & flu relief

acetaminophen 500 mg

dextromethorphan HBr 20 mg

phenylephrine HCl 10 mg

pain reliever/fever reducer

cough suppressant &

nasal decongestant

relieves:

nasal & sinus congestion

cough

body ache

sore throat pain

headache

fever

Green Tea & Honey Lemon flavors

6 PACKETS

nighttime

severe cold cough & flu relief

acetaminophen 650 mg

diphenhydramine HCl 25 mg

phenylephrine HCl 10 mg pain reliever/fever reducer antihistamine/ cough suppressant

& nasal decongestant

relieves:

nasal congestion

cough

runny nose

sneezing

body ache

sore throat pain

headache

fever

honey lemon infused with white tea flavors $% \left(1\right) =\left(1\right) \left(1\right) \left($

6 PACKETS







COMBO PACK

Compare to the active ingredients of Theraflu® Severe Cold and Theraflu® Nighttime Severe Cold & Cough

daytime severe cold cough & flu relief

acetaminophen 500 mg dextromethorphan HBr 20 mg phenylephrine HCI 10 mg

> pain reliever/fever reducer cough suppressant & nasal decongestant

nasal & sinus congestion Honey Lemon flavors sore throat pain headache 6 PACKETS

nighttime severe cold cough & flu relief

acetaminophen 650 mg diphenhydramine HCl 25 mg phenylephrine HCI 10 mg

> pain reliever/fever reducer antihistamine/ cough suppressant & nasal decongestant

nasal congestion cough runny nose sneezing body ache sore throat pain headache fever



honey lemon infused with white tea flavors

6 PACKETS

DO NOT TAKE BOTH PRODUCTS AT THE SAME TIME OR TAKE MORE THAN 5 DOSES IN TOTAL IN ANY 24 HOUR PERIOD.

DO NOT TAKE A DOSE OF THE NIGHTTIME PRODUCT SOONER THAN 4 HOURS AFTER THE LAST DOSE OF MULTI-SYMPTOM PRODUCT UNLESS DIRECTED BY YOUR DOCTOR.

Multi-Symptom Severe Cold

Drug Facts

Active ingredients (in each packet) Purposes Acetaminophen 500 mg... Pain reliever/fever reducer Dextromethorphan HBr 20 mg. .Cough suppressant Phenylephrine HCl 10 mg... ..Nasal decongestant

Uses ■ temporarily relieves these symptoms due to a cold:

- minor aches and pains ■ minor sore throat pain
 ■ headache nasal and sinus congestion cough due to minor throat and bronchial irritation

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

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.
- 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.

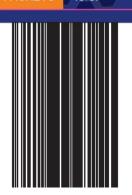
Ask a doctor before use if you have

■ liver disease

- heart disease
 high blood pressure
 thyroid disease ■ trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, or emphysema
- a sodium-restricted diet

Ask a doctor or pharmacist before use if you are taking the blood thinning

When using this product do not exceed recommended dosage





cough suppressant & nasal decongestant antihistamine/ pain reliever/fever reducer

gm 01 IOH ənindqəlvnədq gm 039 **nərdqonims təəs** gm 32 IOH ənimsibydnədqib

Teller condh & flu severe cold amittdgine

Do notuse it printed pack ets are torn or punctured

usasi decongestant con du anb bressant & bain reliever/tever reducer

phenylephrine HCl 10 mg dextromethorphan HBr 20 mg acetaminophen 500 mg

relief condp & flu severe cold daytime

YDAMRAHY **DIA** RITE

Drug Facts (continued)

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

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.

- do not use more than directed (see overdose warning)
 take every 4 hours, while symptoms persist. Do not take more than 6 packets in 24 hours unless directed by a doctor.

Drug Facts (continued)

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adults and children 12 years of age and over	one packet
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Other information

- each packet contains: potassium 10 mg and sodium 25 mg
- phenylketonurics: contains phenylalanine 22 mg per packet ■ store at 20-25°C (68-77°F). Protect product from heat

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

Nighttime Flu & Severe Cold & Cough

Drug Facts

Active ingredients (in each packet) Acetaminophen 650 mg

Phenylephrine HCl 10 mg.

..Pain reliever/fever reducer Diphenhydramine HCl 25 mg.... Antihistamine/cough suppressant

Uses ■ temporarily relieves these symptoms due to a cold: minor aches and pains

- nasal and sinus congestion headache itchy nose or throat runny nose
- minor sore throat pain

...Nasal decongestant

- itchy, watery eyes due to hav fever
- cough due to minor throat and bronchial irritation ■ temporarily reduces fever

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
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Drug Facts (continued)

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

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- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist
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■ thyroid disease Ask a doctor before use if you have

- high blood pressure
- liver disease
 heart 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

Drug Facts (continued)

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 dosag 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

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24 Hours unless uncoted by a doctor.	
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Questions or comments? 1-800-719-9260

*These products are not manufactured or distributed by GSK Consumer Healthcare distributor of Theraflu® Severe Cold and







DISTRIBUTED BY: RITE AID 30 HUNTER LANE CAMP HILL, PA 17011

MADE IN MEXICO





DAYTIME NIGHTTIME SEVERE COLD COUGH AND FLU RELIEF

acetaminophen, dextromethorphan hbr, phenylephrine hcl, diphenhydramine hcl kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:11822-2050

Packaging

l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 N	DC:11822-2050-1	1 in 1 CARTON; Type 0: Not a Combination Product	06/24/2019	

Quantity of Parts

Q cacara	Quantity of Larts		
Part #	Package Quantity	Total Product Quantity	
Part 1	6 PACKET	6	
Part 2	6 PACKET	6	

Part 1 of 2

DAYTIME SEVERE COLD COUGH AND FLU RELIEF

acetaminophen, dextromethorphan hbr, phenylephrine hcl powder, for solution

Product Information

Route of Administration ORAL

Active Ingredient/Active Moiety

retive ingredient/retive morety		
Ingredient Name	Basis of Strength	Strength
ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D)	ACETAMINOPHEN	500 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9 D2RTI9 KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg
PHENYLEPHRINE HYDRO CHLO RIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6 MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg

Inactive 1	lng re die nts
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Ingredient Name	Strength

ACESULFAME POTASSIUM (UNII: 23OV73Q5G9)

ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)	
ASPARTAME (UNII: Z0H242BBR1)	
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
MALTO DEXTRIN (UNII: 7CVR7L4A2D)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SUCROSE (UNII: C151H8 M554)	
TRIBASIC CALCIUM PHO SPHATE (UNII: 91D9 GV0 Z28)	

Product Characteristics		
Color		Score
Shape		Size
Flavor	HONEY (green tea), LEMON (green tea)	Imprint Code
Contains		

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		1 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341		

Part 2 of 2

NIGHTTIME SEVERE COLD COUGH AND FLU RELIEF

acetaminophen, diphenhydramine hcl, phenylephrine hcl powder, for solution

Product Information

Route of Administration ORAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D)	ACETAMINOPHEN	650 mg		
DIPHENHYDRAMINE HYDRO CHLO RIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg		
PHENYLEPHRINE HYDRO CHLO RIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg		

Inactive Ingredients		
Ingredient Name	Strength	
ACESULFAME POTASSIUM (UNII: 23OV73Q5G9)		
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)		
ASPARTAME (UNII: Z0H242BBR1)		
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)		
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		
MALTO DEXTRIN (UNII: 7CVR7L4A2D)		
SODIUM CITRATE (UNII: 1Q73Q2JULR)		
SUCROSE (UNII: C151H8M554)		
TRIBASIC CALCIUM PHO SPHATE (UNII: 91D9GV0Z28)		

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

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		1 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	06/24/2019	

Labeler - Rite Aid Corporation (014578892)

Revised: 8/2019 Rite Aid Corporation