

**COLD AND FLU NIGHTTIME- acetaminophen, dextromethorphan hbr,  
triprolidine hcl solution  
Walgreen Company**

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**Walgreens 44-063**

***Active ingredients (in each 20 mL)***

Acetaminophen 650 mg  
Dextromethorphan HBr 20 mg  
Triprolidine HCl 2.5 mg

***Purpose***

Pain reliever/fever reducer  
Cough suppressant  
Antihistamine

***Uses***

- temporarily relieves these common cold and flu symptoms:
  - cough
  - headache
  - runny nose
  - sneezing
  - sore throat
  - itching of the nose or throat
  - minor aches and pains
  - itchy, watery eyes due to hay fever
- temporarily reduces fever
- controls cough to help you get to sleep

***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:

- rash
- blisters
- skin reddening

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**

- 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.
- if you have ever had an allergic reaction to this product or any of its ingredients

**Ask a doctor before use if you have**

- cough that occurs with too much phlegm (mucus)
- difficulty in urination due to enlargement of the prostate gland
- liver disease
- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- persistent or chronic cough 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**
- excitability may occur, especially in children
- avoid alcoholic beverages
- marked drowsiness may occur
- alcohol, sedatives, and tranquilizers may increase drowsiness
- use caution when driving a motor vehicle or operating machinery

**Stop use and ask a doctor if**

- pain or cough gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- new symptoms occur
- redness or swelling is present
- 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 take more than directed**
- do not take more than 4 doses in any 24-hour period
- mL = milliliter
- only use the dose cup provided
- adults and children 12 years and over: 20 mL in dosing cup provided every 4 hours
- children under 12 years: do not use

### ***Other information***

- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- use by expiration date on package

### ***Inactive ingredients***

anhydrous citric acid, FD&C blue #1, FD&C red #40, FD&C yellow #6, flavors, glycerin, propylene glycol, purified water, sodium benzoate, sucralose, xanthan gum

### ***Questions or comments?***

**1-800-426-9391**

### ***Principal display panel***

NDC 0363-8063-45

### ***Walgreens***

WALGREENS PHARMACIST RECOMMENDED†

Compare to the active ingredients  
in Mucinex® NIGHTSHIFT® Cold & Flu††

### **NIGHTTIME**

#### **Cold & Flu**

**ACETAMINOPHEN / PAIN RELIEVER / FEVER REDUCER**

**DEXTROMETHORPHAN HBr / COUGH SUPPRESSANT**

**TRIPROLIDINE HCl / ANTIHISTAMINE**

Multi-Symptom

- Relieves cough, fever, sore throat,  
runny nose & sneezing
- 12 years & older

**6 FL OZ (177 mL)**

**TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY  
SEAL UNDER CAP IS BROKEN OR MISSING**

**PARENTS:**

**Learn about teen medicine abuse  
[www.StopMedicineAbuse.org](http://www.StopMedicineAbuse.org)**

50844 ORG042306345

†Our pharmacists  
recommend the  
Walgreens brand.

We invite you to  
compare to  
national brands.

††This product is not manufactured or  
distributed by RB Health (US) LLC, owner  
of the registered trademark Mucinex®  
NIGHTSHIFT® Cold & Flu.

DISTRIBUTED BY: **WALGREEN CO.**  
**DEERFIELD, IL 60015**

**100% SATISFACTION GUARANTEED**

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Our pharmacists recommend the Walgreens brand. We invite you to compare to national brands.

**PARENTS:** Learn about teen medicine abuse [www.StopMedicineAbuse.org](http://www.StopMedicineAbuse.org)

PEEL BACK TAB TO READ COMPLETE DRUG FACTS AND INFORMATION

ITEM 892389 W00000-0000-0  
B-063-45 ORG  
ORG042306345

11917 03710 3  
W3ORG0923-F

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**Drug Facts** **TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING**

**Active ingredients (in each 20 mL)** **Purpose**  
 Acetaminophen 650 mg ..... Pain reliever/fever reducer  
 Dextromethorphan HBr 20 mg ..... Cough suppressant  
 Triprolidine HCl 2.5 mg ..... Antihistamine

**Uses** ■ temporarily relieves these common cold and flu symptoms: ■ cough ■ headache ■ runny nose ■ sneezing ■ sore throat ■ itching of the nose or throat ■ minor aches and pains ■ itchy, watery eyes due to hay fever ■ temporarily reduces fever ■ controls cough to help you get to sleep

**Warnings**  
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**Allergy alert:** Acetaminophen may cause severe skin reactions. Symptoms may include: ■ rash ■ blisters

No Print / No Varnish Area  
Lot # and Exp. Info

**Drug Facts (continued)**

■ skin reddening  
 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** ■ 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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**Drug Facts (continued)**

drowsiness ■ avoid alcoholic beverages  
 ■ use caution when driving a motor vehicle or operating machinery™

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**Questions or comments?** 1-800-426-9391

Walgreens 44-063

## COLD AND FLU NIGHTTIME

acetaminophen, dextromethorphan hbr, triprolidine hcl solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0363-8063
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg in 20 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 20 mL
TRIPROLIDINE HYDROCHLORIDE (UNII: YAN7R5L890) (TRIPROLIDINE - UNII:2L8T9S52QM)	TRIPROLIDINE HYDROCHLORIDE	2.5 mg in 20 mL

## Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

## Product Characteristics

Color	blue	Score	
Shape		Size	
Flavor	FRUIT	Imprint Code	
Contains			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-8063-45	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/23/2023	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	10/23/2023	

**Labeler** - Walgreen Company (008965063)

## Establishment

Name	Address	ID/FEI	Business Operations
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LNK International, Inc.

967626305

manufacture(0363-8063) , pack(0363-8063)

Revised: 10/2023

Walgreen Company