

**COLD AND FLU SEVERE, DAYTIME, NIGHTTIME- acetaminophen,  
chlorpheniramine maleate, dextromethorphan hbr, guaifenesin,  
phenylephrine hcl  
Rite Aid Corporation**

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**Rite Aid 44-503A473C**

***Active ingredients (in each caplet) (Daytime Cold & Flu Severe)***

Acetaminophen 325 mg  
Dextromethorphan HBr 10 mg  
Guaifenesin 200 mg  
Phenylephrine HCl 5 mg

***Purpose***

Pain reliever/fever reducer  
Cough suppressant  
Expectorant  
Nasal decongestant

***Active ingredients (in each caplet) (Nighttime Cold & Flu Severe)***

Acetaminophen 325 mg  
Chlorpheniramine maleate 2 mg  
Dextromethorphan HBr 10 mg  
Phenylephrine HCl 5 mg

***Purpose***

Pain reliever/fever reducer  
Antihistamine  
Cough suppressant  
Nasal decongestant

***Uses***

- temporarily relieves these common cold and flu symptoms:
  - cough
  - headache
  - sore throat
  - nasal congestion
  - minor aches and pains
  - sinus congestion and pressure
  - sneezing and runny nose (***Nighttime only***)
- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive (***Daytime only***)

- helps clear nasal passages
- relieves cough to help you sleep
- 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**

- 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**

- high blood pressure
- liver disease
- diabetes
- thyroid disease
- heart disease
- glaucoma (**Nighttime only**)
- cough that occurs with too much phlegm (mucus)
- difficulty in urination due to enlargement of the prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- a breathing problem such as emphysema or chronic bronchitis (**Nighttime only**)

## **Ask a doctor or pharmacist before use if you are**

- taking the blood thinning drug warfarin

- taking sedatives or tranquilizers (**Nighttime only**)

### **When using this product**

- **do not exceed recommended dosage**
- excitability may occur, especially in children (**Nighttime only**)
- marked drowsiness may occur (**Nighttime only**)
- avoid alcoholic beverages (**Nighttime only**)
- use caution when driving a motor vehicle or operating machinery (**Nighttime only**)
- alcohol, sedatives, and tranquilizers may increase drowsiness (**Nighttime only**)

### **Stop use and ask a doctor if**

- nervousness, dizziness, or sleeplessness occur
- pain, nasal congestion, or cough gets worse or lasts more than 7 days
- new symptoms occur
- fever gets worse or lasts more than 3 days
- 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.

### **Do not take DAYTIME and NIGHTTIME products at the same time.**

### **Directions**

- **do not take more than directed**
- adults and children 12 years and over
  - take 2 caplets every 4 hours
  - swallow whole – do not crush, chew, or dissolve
  - do not take more than 10 caplets in 24 hours
- children under 12 years: ask a doctor

### **Other information**

- **each caplet contains:** sodium 3 mg
- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

### **Inactive ingredients (Daytime only)**

corn starch, crospovidone, D&C yellow #10 aluminum lake, flavor, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, sucralose, talc, titanium dioxide

***Inactive ingredients (Nighttime only)***

corn starch, crospovidone, FD&C blue #1 aluminum lake, FD&C blue #2 aluminum lake, flavor, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, sucralose, talc, titanium dioxide

***Questions or comments?***

**1-800-426-9391**

***Principal display panel***

NDC 11822-5034-8

Compare to the active ingredients in

**Tylenol® COLD + FLU SEVERE Day & Tylenol® COLD + FLU SEVERE Night\***

**COLD & FLU SEVERE**

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**DAYTIME**

**ACETAMINOPHEN**

DEXTROMETHORPHAN HBr

GUAIFENESIN

PHENYLEPHRINE HCl

PAIN RELIEVER/FEVER REDUCER

COUGH SUPPRESSANT • EXPECTORANT

NASAL DECONGESTANT

Fever, Headache, Sore Throat,

Nasal Congestion, Cough,

Mucus, Chest Congestion

ACTUAL SIZE

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**NIGHTTIME**

**ACETAMINOPHEN**

CHLORPHENIRAMINE MALEATE

DEXTROMETHORPHAN HBr

PHENYLEPHRINE HCl

PAIN RELIEVER/FEVER REDUCER

ANTIHISTAMINE • COUGH SUPPRESSANT

NASAL DECONGESTANT

Fever, Headache, Sore Throat,

Runny Nose, Cough,

Nasal Congestion

ACTUAL SIZE

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PSEUDOEPHEDRINE FREE

**16**

**DAYTIME**

CAPLETS

**8**

**NIGHTTIME**

CAPLETS

**TAMPER EVIDENT: DO NOT USE IF  
PACKAGE IS OPENED OR IF BLISTER  
UNIT IS TORN, BROKEN OR SHOWS**

## **ANY SIGNS OF TAMPERING**

\*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Tylenol® COLD + FLU SEVERE Day & Tylenol® COLD + FLU SEVERE Night.

## **SATISFACTION GUARANTEE**

If you're not satisfied, we'll happily refund your money.

**DISTRIBUTED BY:** RITE AID,  
200 NEWBERRY COMMONS  
ETTERS, PA 17319 [www.riteaid.com](http://www.riteaid.com)

50844 REV0922A50347308

## **PARENTS:**

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

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

**Drug Facts**

**Active ingredients (in each caplet) (Daytime Cold & Flu Severe)**

Acetaminophen 325 mg	Pain reliever/fever reducer
Dextromethorphan HBr 10 mg	Cough suppressant
Guaifenesin 200 mg	Expectorant
Phenylephrine HCl 5 mg	Nasal decongestant

**Purpose**

**Active ingredients (in each caplet) (Nighttime Cold & Flu Severe)**

Acetaminophen 325 mg	Pain reliever/fever reducer
Chlorpheniramine maleate 2 mg	Antihistamine
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Phenylephrine HCl 5 mg	Nasal decongestant

**Purpose**

**Uses**

- temporarily relieves these common cold and flu symptoms:
  - cough
  - headache
  - sore throat
  - nasal congestion
  - minor aches and pains
  - sinus congestion and pressure
  - sneezing and runny nose (Nighttime only)

**Drug Facts (continued)**

- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive (Daytime only)
- helps clear nasal passages
- relieves cough to help you sleep
- 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
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- Allergy alert:** Acetaminophen may cause severe skin reactions. Symptoms may include:
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  - rash
 If a skin reaction occurs, stop use and seek medical help right away.
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**Do not use**

- with any other drug containing acetaminophen (prescription or

TYPE SAFETY

PEEL HERE FOR MORE DRUG FACTS

**COLD & FLU SEVERE**



**DAYTIME**



**NIGHTTIME**

B-1702503A473008R3  
REV0922A50347308

NDC 11822-5034-8

Compare to the active ingredients in

Tylenol® COLD + FLU SEVERE Day & Tylenol® COLD + FLU SEVERE Night\*

**COLD & FLU SEVERE**



**DAYTIME**



**NIGHTTIME**

**ACETAMINOPHEN  
DXTROMETHORPHAN HBr  
GUAIFENESIN  
PHENYLEPHRINE HCl**

PAIN RELIEVER/FEVER REDUCER  
COUGH SUPPRESSANT • EXPECTORANT  
NASAL DECONGESTANT

Fever, Headache, Sore Throat,  
Nasal Congestion, Cough,  
Mucus, Chest Congestion

ACTUAL SIZE

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CHLORPHENIRAMINE MALEATE  
DXTROMETHORPHAN HBr  
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PAIN RELIEVER/FEVER REDUCER  
ANTIHISTAMINE • COUGH SUPPRESSANT  
NASAL DECONGESTANT

Fever, Headache, Sore Throat,  
Runny Nose, Cough,  
Nasal Congestion

ACTUAL SIZE

NO PRINT AREA  
LOT & EXP. DATE

TAMPER EVIDENT: DO NOT USE IF  
PACKAGE IS OPENED OR IF BLISTERS  
OR BUBBLES ARE VISIBLE OR  
IF THERE ARE CRACKS OR  
SMALL HOLES IN THE SEAL



PSEUDOEPHEDRINE FREE

**16**  
DAYTIME  
CAPLETS

**8**  
NIGHTTIME  
CAPLETS

**COLD & FLU SEVERE**  
DAYTIME NIGHTTIME

www.StopAches.com  
**PARENTS:**  
LEARN HOW BEST MEDICINE WORKS

DISTRIBUTED BY: RTE PHARM,  
200 NEUMBERY COMMONS  
LETTERS, PA 17319 www.rteaid.com

**SATISFACTION  
GUARANTEE**  
If you're not satisfied, we'll  
happily refund your money.

50844  
REV0922A50347308

50844

Tylenol® COLD + FLU SEVERE Day & Tylenol® COLD + FLU SEVERE Night

\*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark

**Drug Facts (continued)**

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.
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- swallow whole - do not crush, chew, or dissolve
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**Other information**

- each caplet contains: sodium 3 mg
- TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

**Drug Facts (continued)**

**Inactive ingredients (Daytime only)**

corn starch, croscopolone, D&C yellow #10 aluminum lake, flavor, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, sucralose, talc, titanium dioxide

**Inactive ingredients (Nighttime only)**

corn starch, croscopolone, FD&C blue #1 aluminum lake, FD&C blue #2 aluminum lake, flavor, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, sucralose, talc, titanium dioxide

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

50844  
REV0922A50347308  
R-1702503A473008DF

TYPE SAFETY

Rite Aid 44-503A473C

## COLD AND FLU SEVERE, DAYTIME, NIGHTTIME

acetaminophen, chlorpheniramine maleate, dextromethorphan hbr, guaifenesin, phenylephrine hcl kit

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11822-5034
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### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-5034-8	1 in 1 CARTON; Type 0: Not a Combination Product	06/30/2021	

### Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	2 BLISTER PACK	16
Part 2	1 BLISTER PACK	8

### Part 1 of 2

## COLD AND FLU SEVERE DAYTIME

acetaminophen, dextromethorphan hbr, guaifenesin phenylephrine hcl tablet, film coated

### Product Information

<b>Item Code (Source)</b>	NDC:11822-5347
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<b>Route of Administration</b>	ORAL
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### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>CROSPVIDONE, UNSPECIFIED</b> (UNII: 2S7830E561)	

<b>D&amp;C YELLOW NO. 10 ALUMINUM LAKE</b> (UNII: CQ3XH3DET6)
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)
<b>POLYVINYL ALCOHOL, UNSPECIFIED</b> (UNII: 532B59J990)
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)
<b>SODIUM STARCH GLYCOLATE TYPE A POTATO</b> (UNII: 5856J3G2A2)
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)
<b>TALC</b> (UNII: 7SEV7J4R1U)
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)

### Product Characteristics

<b>Color</b>	yellow	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	19mm
<b>Flavor</b>	MENTHOL	<b>Imprint Code</b>	44;503
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-5347-9	8 in 1 BLISTER PACK; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	06/30/2021	

## Part 2 of 2

### COLD AND FLU SEVERE NIGHTTIME

acetaminophen, chlorpheniramine maleate, dextromethorphan hbr, phenylephrine hcl tablet, film coated

### Product Information

<b>Item Code (Source)</b>	NDC:11822-4753
<b>Route of Administration</b>	ORAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
<b>CHLORPHENIRAMINE MALEATE</b> (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	2 mg
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

## Inactive Ingredients

Ingredient Name	Strength
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>CROSPVIDONE, UNSPECIFIED</b> (UNII: 2S7830E561)	
<b>FD&amp;C BLUE NO. 1 ALUMINUM LAKE</b> (UNII: J9EQA3S2JM)	
<b>FD&amp;C BLUE NO. 2 ALUMINUM LAKE</b> (UNII: 4AQJ3LG584)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POLYVINYL ALCOHOL, UNSPECIFIED</b> (UNII: 532B59J990)	
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>SODIUM STARCH GLYCOLATE TYPE A POTATO</b> (UNII: 5856J3G2A2)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

## Product Characteristics

<b>Color</b>	blue	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	17mm
<b>Flavor</b>	MENTHOL	<b>Imprint Code</b>	44;473
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-4753-9	8 in 1 BLISTER PACK; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	06/30/2021	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	06/30/2021	

**Labeler** - Rite Aid Corporation (014578892)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(11822-5034) , pack(11822-5034)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(11822-5034)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(11822-5034)

Revised: 5/2025

Rite Aid Corporation