

**COLD AND FLU SEVERE, DAY, NIGHT- acetaminophen, chlorpheniramine maleate, dextromethorphan hbr. guaifenesin, phenylephrine hcl**  
**L.N.K. International, Inc.**

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**Quality Plus 44-503A473C-08**

***Active ingredients (in each caplet) (daytime)***

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)***

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.

## **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***

**QUALITY**

**+PLUS**

NDC 50844-547-08

\*Compare to active ingredients in  
Tylenol® COLD + FLU  
SEVERE DAY & NIGHT

**SEVERE**

**COLD & FLU**

**DAY**

**Acetaminophen,**

Dextromethorphan HBr,

Guaifenesin,

Phenylephrine HCl

PAIN RELIEVER/

FEVER REDUCER,

COUGH SUPPRESSANT,

EXPECTORANT,

NASAL DECONGESTANT

ACTUAL

SIZE

**16**

**Caplets**

**NIGHT**

**Acetaminophen,**

Chlorpheniramine maleate,

Dextromethorphan HBr,

Phenylephrine HCl

PAIN RELIEVER/

FEVER REDUCER,

ANTIHISTAMINE,  
COUGH SUPPRESSANT,  
NASAL DECONGESTANT

ACTUAL  
SIZE

**8**  
**Caplets**

**24 Total 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  
Kenvue Inc., owner of the registered trademark  
Tylenol® COLD + FLU SEVERE DAY & NIGHT.  
50844                      REV0825B50347308

**Do Not Take Daytime and  
Nighttime Products at the  
Same Time.**

Distributed by  
**LNK INTERNATIONAL, INC.**  
60 Arkay Drive  
Hauppauge, NY 11788  
USA

**QUALITY PLUS**

**SEVERE COLD & FLU DAY & NIGHT**

NDC 50844-547-08

\*Compare to active ingredients in Tylenol® COLD + FLU SEVERE DAY & NIGHT

**QUALITY PLUS**

**SEVERE COLD & FLU DAY & NIGHT**

**DAY**

**Acetaminophen, Dextromethorphan HBr, Guaifenesin, Phenylephrine HCl**

PAIN RELIEVER/ FEVER REDUCER, COUGH SUPPRESSANT, EXPECTORANT, NASAL DECONGESTANT

**16 Caplets**

**NIGHT**

**Acetaminophen, Chlorpheniramine maleate, Dextromethorphan HBr, Phenylephrine HCl**

PAIN RELIEVER/ FEVER REDUCER, ANTIHISTAMINE, COUGH SUPPRESSANT, NASAL DECONGESTANT

**8 Caplets**

**24 Total Caplets**



3 50844 54708 8

No print/No varnish area  
Lot no/Exp date

**QUALITY PLUS**

**SEVERE COLD & FLU DAY & NIGHT**

**KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION**

<p><b>Drug Facts</b></p> <p><b>Active ingredients (in each caplet) (daytime)</b></p> <p>Acetaminophen 325 mg Dextromethorphan HBr 10 mg Guaifenesin 200 mg Phenylephrine HCl 5 mg</p> <p><b>Active ingredients (in each caplet) (nighttime)</b></p> <p>Acetaminophen 325 mg Chlorpheniramine maleate 2 mg Dextromethorphan HBr 10 mg Phenylephrine HCl 5 mg</p> <p><b>Uses</b></p> <ul style="list-style-type: none"> <li>■ minor aches and pains</li> <li>■ headache</li> <li>■ nasal congestion</li> <li>■ sinus congestion and pressure</li> <li>■ sneezing and runny nose (nighttime only)</li> <li>■ helps relieve rhinitis (nasal) and the bronchial secretions to make coughs more productive (daytime only)</li> </ul>	<p><b>Drug Facts (continued)</b></p> <p>■ helps clear nasal passages</p> <p>■ relieves cough to help you sleep</p> <p>■ temporarily reduces fever</p> <p><b>Warnings</b></p> <p>Liver warnings: This product contains acetaminophen. Severe liver damage may occur if you take</p> <ul style="list-style-type: none"> <li>■ more than 4,000 mg of acetaminophen in 24 hours</li> <li>■ with other drugs containing acetaminophen</li> <li>■ 3 or more alcoholic drinks every day while using this product</li> </ul> <p><b>Allergy alert:</b> Ketaminophen may cause severe skin reactions.</p> <p>Symptoms may include:</p> <ul style="list-style-type: none"> <li>■ skin redness</li> <li>■ hives</li> <li>■ rash</li> </ul> <p>See the back of this caplet for more information. If you are allergic to any of the ingredients, this product is contraindicated for you. If you are unsure if you are allergic, ask your doctor. If you have had an allergic reaction to any other drug, is contraindicated or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.</p> <p><b>Do not use</b></p> <ul style="list-style-type: none"> <li>■ with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.</li> </ul>
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PEEL HERE FOR MORE DRUG FACTS

B-1603-503A473C-08  
REV0825B50347308

<p><b>Drug Facts (continued)</b></p> <ul style="list-style-type: none"> <li>■ 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.</li> </ul> <p><b>Ask a doctor before use if you have</b></p> <ul style="list-style-type: none"> <li>■ liver disease</li> <li>■ heart disease</li> <li>■ high blood pressure</li> <li>■ thyroid disease</li> <li>■ diabetes</li> <li>■ difficulty in urination due to enlargement of the prostate gland</li> <li>■ cough that occurs with too much phlegm (mucus)</li> <li>■ persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema</li> <li>■ a breathing problem such as emphysema or chronic bronchitis (nighttime only)</li> <li>■ glaucoma (nighttime only)</li> </ul> <p><b>Ask a doctor or pharmacist before use if you are</b></p> <ul style="list-style-type: none"> <li>■ taking the blood thinning drug warfarin</li> <li>■ taking sedatives or tranquilizers (nighttime only)</li> </ul>	<p><b>Drug Facts (continued)</b></p> <p><b>When using this product</b></p> <ul style="list-style-type: none"> <li>■ do not exceed recommended dosage</li> <li>■ excitability may occur, especially in children (nighttime only)</li> <li>■ marked drowsiness may occur (nighttime only)</li> <li>■ alcohol, sedatives, and tranquilizers may increase drowsiness (nighttime only)</li> <li>■ avoid alcoholic beverages (nighttime only)</li> <li>■ use caution when driving a motor vehicle or operating machinery (nighttime only)</li> </ul> <p><b>Stop use and ask a doctor if</b></p> <ul style="list-style-type: none"> <li>■ nervousness, dizziness, or sleeplessness occur</li> <li>■ pain, nasal congestion, or cough gets worse or lasts more than 7 days</li> <li>■ fever gets worse or lasts more than 3 days</li> <li>■ redness or swelling is present</li> <li>■ new symptoms occur</li> <li>■ cough comes back or occurs with rash or headache that lasts. These could be signs of a serious condition.</li> </ul> <p>If pregnant or breast-feeding, ask a health professional before use.</p>
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**Drug Facts (continued)**

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

**Drug Facts (continued)**

**Inactive ingredients (daytime only)**

corn starch, croscopolone, FD&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  
REV0825B50347308  
R-1603-503A473C08-DF

**Quality plus 44-503A473C**

**COLD AND FLU SEVERE, DAY, NIGHT**  
acetaminophen, chlorpheniramine maleate, dextromethorphan hbr. guaifenesin, phenylephrine hcl kit

<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:50844-547

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50844-547-08	1 in 1 CARTON; Type 0: Not a Combination Product	07/15/2021	

## 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 DAY

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

## Product Information

Item Code (Source) NDC:50844-980

Route of Administration ORAL

## Active Ingredient/Active Moiety

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

## Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
CROSPVIDONE, UNSPECIFIED (UNII: 2S7830E561)	
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
SILICON DIOXIDE (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		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	07/15/2021	

## Part 2 of 2

### COLD AND FLU SEVERE NIGHT

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

### Product Information

<b>Item Code (Source)</b>	NDC:50844-981
<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>CROSPROVIDONE, 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		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	07/15/2021	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/15/2021	

**Labeler** - L.N.K. International, Inc. (038154464)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		832867837	manufacture(50844-547) , pack(50844-547)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		832867894	manufacture(50844-547)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		117025878	manufacture(50844-547)

Revised: 2/2026

L.N.K. International, Inc.