

**COMPLETE COLD RELIEF- acetaminophen, dextromethorphan hbr,  
guaifenesin, phenylephrine hcl solution  
Belmora LLC**

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**Buckleys 44-005**

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

Acetaminophen 650 mg  
Dextromethorphan HBr 20 mg  
Guaifenesin 400 mg  
Phenylephrine HCl 10 mg

***Purpose***

Pain reliever/fever reducer  
Cough suppressant  
Expectorant  
Nasal decongestant

***Uses***

- temporarily relieves these common cold and flu symptoms:
  - minor aches and pains
  - sinus congestion and pressure
  - stuffy nose
  - nasal congestion
  - sore throat
  - cough
  - headache
- temporarily promotes nasal and/or sinus drainage
- temporarily reduces fever
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

***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:

- blisters
- rash

- 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)
- liver disease
- thyroid disease
- heart disease
- difficulty in urination due to enlargement of the prostate gland
- high blood pressure
- diabetes
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema

### **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**

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

### **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 6 doses in any 24-hour period
- mL = milliliter
- only use the dose cup provided
- dose as follows or as directed by a doctor
- 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***

- **each 20 mL contains:** sodium 10 mg
- 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, flavors, glycerin, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate dihydrate, sodium metabisulfite, sorbitol, sucralose

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

**1-888-779-2877 M-F 9AM-5PM EST**

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

**Belmora LLC**

NDC 27854-299-00

**MAXIMUM STRENGTH**

**Buckleys**

**COMPLETE COLD RELIEF**

**ACETAMINOPHEN -**

PAIN RELIEVER/FEVER REDUCER

DEXTROMETHORPHAN HBr - COUGH SUPPRESSANT

GUAIFENESIN - EXPECTORANT

PHENYLEPHRINE HCl - NASAL DECONGESTANT

RELIEVES:

✓ HEADACHE, FEVER & SORE THROAT

✓ SINUS PRESSURE AND CONGESTION

✓CONTROLS COUGH, THINS & LOOSENS MUCUS

**AGES 12 & OVER**

F-005-45 REV A

**6 FL OZ (177 mL)**

MIXED BERRY FLAVORED

**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)**

**Distributed by: Belmora LLC, 3033 Wilson Blvd. Suite 700**

**Arlington, VA 22201**

50844 REV0523A00545



NDC 27854-299-00

MAXIMUM STRENGTH

# Buckleys

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**PARENTS:**  
Learn about how medicine works  
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**PEEL BACK TAB TO READ COMPLETE  
DRUG FACTS AND INFORMATION**

B-005-45 REV A

## Drug Facts

**TAMPER EVIDENT: DO NOT USE IF IMPRINTED  
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Acetaminophen 650 mg.....	Pain reliever/fever reducer
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**Uses** ■ temporarily relieves these common cold and flu symptoms: ■ minor aches and pains ■ sinus congestion and pressure ■ stuffy nose ■ nasal congestion ■ sore throat ■ cough ■ headache ■ temporarily promotes nasal and/or sinus drainage ■ temporarily reduces fever ■ helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

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

## Drug Facts (continued)

- 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: ■ blisters ■ rash ■ skin reddening. If a skin reaction occurs, stop use and seek medical help right away.

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- cough that occurs with too much phlegm (mucus)

HINGE

## Drug Facts (continued)

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HINGE

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 ■ difficulty in urination due to enlargement of the prostate gland ■ high blood pressure ■ diabetes  
 ■ persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema

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permitted between 15°-30°C (59°-86°F)  
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**Inactive ingredients** anhydrous citric acid, FD&C blue #1, FD&C red #40, flavors, glycerin, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate dihydrate, sodium metabisulfite, sorbitol, sucralose

**Questions or comments?**  
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### Buckleys 44-005

## COMPLETE COLD RELIEF

acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:27854-299
<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	650 mg in 20 mL
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 20 mL
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 20 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>TRISODIUM CITRATE DIHYDRATE</b> (UNII: B22547B95K)	
<b>SODIUM METABISULFITE</b> (UNII: 4VON5FNS3C)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	

### Product Characteristics

<b>Color</b>	blue	<b>Score</b>	
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<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	BERRY	<b>Imprint Code</b>	
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:27854-299-00	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/23/2022	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	02/23/2022	

**Labeler** - Belmora LLC (112753244)