

**CODEINE-GUAIFENESIN- codeine phosphate and guaifenesin solution**  
**NuCare Pharmaceuticals, Inc.**

*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.*

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**Codeine-Guaifenesin Oral Solution**

***Active ingredient***

***(in each teaspoonful (5 mL))***

Codeine Phosphate USP 10 mg

***Purpose***

Antitussive

***Active ingredient***

***(in each teaspoonful (5 mL))***

Guaifenesin USP 100 mg

***Purpose***

Expectorant

***Uses***

- temporarily relieves:
- cough due to minor throat and bronchial irritation as may occur with a cold or inhaled irritants
- your cough to help you sleep
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and makes cough more productive.

***Warnings***

**Ask your doctor before use if**

- you have a persistent cough, this may be a sign of a serious condition
- you have a persistent cough such as occurs with smoking, asthma, chronic bronchitis or emphysema
- you have a cough that is accompanied by excessive phlegm (mucus)
- you have chronic pulmonary disease or shortness of breath
- giving to a child who is taking other drugs

**When using this product**

- giving a higher dose than recommended by a doctor could result in serious side effects for your child. A special measuring device should be used to give an accurate dose of this product to children under 6 years of age.
- may cause or aggravate constipation

**Stop use and ask a doctor if**

- symptoms do not improve within 7 days, tend to recur or are accompanied by fever and rash or persistent headache. These may be symptoms 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.

***Directions***

- do not exceed 6 doses in 24 hours.

Adults and children 12 years of age and over:	2 tsp (10 mL) every 4 hours, or as directed by a doctor.
Children 6 to under 12 years of age:	1 tsp (5 mL) every 4 hours, or as directed by a doctor.
Children under 6 years of age:	Consult a doctor.

***Other information***

Store at controlled room temperature 15°-30°C (59°-86°F).

***Inactive ingredients***

Cherry Flavor, Citric Acid, Glycerin, Propylene glycol, Purified Water, Sodium Citrate, Sodium Saccharin, Sorbitol.

**Principal Display Panel**

# NuCare Pharmaceuticals, Inc.

NDC: 68071-5173-4  
Cod.Phos.10mg/Guaifenesin 100mg/5mL

Cod.Phos.10mg/Guaifenesin 100mg/5mL  
Lot: 000000 NDC: 68071-5173-04  
MFR NDC: 58657-500 04 Exp.: 00-00  
Serial# 00000000002

Cod.Phos.10mg/Guaifenesin 100mg/5mL  
Lot: 000000 NDC: 68071-5173-04  
MFR NDC: 58657-500 04 Exp.: 00-00  
Serial# 00000000002

4oz Oral Soln.

See manufacturer's label  
for full list of ingredients.

Product #: R0274004  
Rx Only



GTIN 00368071517347  
Serial# 00000000002  
Exp. Date 00-00  
LOT#: 000000

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

WARNING: KEEP OUT OF REACH OF CHILDREN

STORE AT CONTROLLED TEMPERATURE 59-86°F.

## CODEINE-GUAIFENESIN

codeine phosphate and guaifenesin solution

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-5173(NDC:58657-500)
Route of Administration	ORAL	DEA Schedule	CV

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CODEINE PHOSPHATE (UNII: GSL05Y1MN6) (CODEINE ANHYDROUS - UNII:UX6OWY2V7J)	CODEINE PHOSPHATE	10 mg in 5 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 5 mL

### Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL (UNII: 506T60A25R)	

### Product Characteristics

Color		Score	
Shape		Size	

<b>Flavor</b>	CHERRY	<b>Imprint Code</b>	
<b>Contains</b>			
<b>Packaging</b>			
<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>
<b>1</b>	NDC:68071-5173-4	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2020
<b>Marketing Information</b>			
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC monograph final	part341	04/01/2014	

**Labeler** - NuCare Pharmaceuticals,Inc. (010632300)

Establishment			
Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals,Inc.		010632300	relabel(68071-5173)