

CODEINE-GUAIFENESIN- codeine phosphate and guaifenesin solution
ATLANTIC BIOLOGICALS CORP.

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.

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, Polyethylene Glycol, Purified Water, Sodium Citrate, Sodium Saccharin, Sorbitol.

Principal Display Panel

NDC 17856-0500-5

Codeine-Guaifenesin Oral Solution

10-100 mg/5 mL

Antitussive

Expectorant

16 fl. oz. (473 mL)

17856-0500-05
CODEINE- GUAIFENESIN
ORAL SOLUTION 10 MG/100
MG/ 5 ML SUGAR FREE,
ALCOHOL FREE, DYE FREE



See package insert for indications and dosage schedule

Store at controlled room temperature 15°-30°C
(59°-86° F).**** Keep this and all Medication
out of the reach of children



17856-0500-05

Dosage: 5 ML

CODEINE-GUAIFENESIN

Qty: 72 CUPS



GTIN: 00117856050056

S/N: 01264401

Exp: 07/08/21

Lot: 012644

CV

Packaged by: Unit Dose Solutions
Morrisville, NC 27560

Distributed by: Atlantic Biologicals Corp.
Miami FL 33179

Rev.09/19

Call to Reorder: 800.509.7592

CODEINE-GUAIFENESIN

codeine phosphate and guaifenesin solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:17856-0500(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		
Flavor	CHERRY	Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17856-0500-5	5 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product	01/26/2021	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC MONOGRAPH FINAL	part341	04/01/2014		

Labeler - ATLANTIC BIOLOGICALS CORP. (047437707)

Registrant - ATLANTIC BIOLOGICALS CORP. (047437707)

Establishment			
Name	Address	ID/FEI	Business Operations
ATLANTIC BIOLOGICALS CORP.		047437707	repack(17856-0500)