BACTIMICINA COUGH AND COLD- dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride solution
DLC Laboratories, 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.

BACTIMICINA ® COUGH AND COLD

Drug Facts

Active ingredients (in each 5 mL = 1 teaspoon)	***Purposes***
Dextromethorphan HBr, USP 10 mg	Cough Suppressant
Guaifenesin, USP 100 mg	Expectorant
Phenylephrine HCl, USP 5 mg	Nasal Decongestant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes
- temporarily relieves these symptoms occurring with a cold:
  - nasal congestion
  - cough due to minor throat and bronchial irritation

Warnings

Do not use

- 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

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema

Ask a doctor or pharmacist before use if you are taking any other oral nasal decongestant or stimulant.

When using this product do not use more than directed

Stop use and ask a doctor if

- you get nervous, dizzy, or sleepless
- symptoms do not get better within 7 days or are accompanied by fever
- cough lasts more than 7 days, comes back, or is accompanied by fever, rash, or persistent headache. 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.

Directions

- do not take more than 6 doses in any 24-hour period
- this adult product is not intended for children under 12 years old
- mL=mililiter;tsp=teaspoonful

Age (years)	Dose
12 and over	2 teaspoons (10 mL) every 4 hours
Under 12	do not use

Other information

- **each teaspoon (5 mL) contains:** sodium 2 mg
- store at 15-30°C (59-86°F)
- measure only with dosage cup provided

Inactive ingredients

citric acid, FD&C red no. 40, flavor, glycerin, propylene glycol, purified water, saccharin sodium, sodium benzoate, sorbitol

Questions

1-800-858-3889

Manufactured by
DLC Laboratories, Inc.
Paramount, CA 90723 USA

Principal Display

Triple Action Voltee Para Esponol

TRUSTED SINCE 1978

New Look

Adult Formula

Bactimicina

Ages 12+ Years

Multi-Symptom • Liquid

Cough & Cold

Dextromethorphan HBr (Cough Suppresant)

Guaifenesin (Expectorant)

Phenylephrine HCl (Nasal Decongestant)

Non-Drowsy

Cough • Stuffy Nose

Chest Congestion • Mucus

Alcohol Free

Tussin CF

Natural Stawberry Flavor

4 FL OZ (118 mL)

BACTIMICINA COUGH AND COLD

dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24286-1546
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH)	DEXTROMETHORPHAN	10 mg

(DEXTROMETHORPHAN - UNII:7355X3ROTS)	HYDROBROMIDE	in 5 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 5 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL (UNII: 506T60A25R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	STRAWBERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24286-1546-4	1 in 1 BOX	11/06/2013	
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	11/06/2013	

Labeler - DLC Laboratories, Inc. (093351930)

Establishment

Name	Address	ID/FEI	Business Operations
DLC Laboratories, Inc.		093351930	manufacture(24286-1546) , label(24286-1546)