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

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 Drug	M012	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)

Revised: 10/2024 DLC Laboratories, Inc.