BACTIMICINA CHILDRENS COUGH AND COLD- chlorpheniramine maleate, dextromethorphan hydrobromide liquid 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 Liquid Children's Cough and Cold

Drug Facts

Active ingredients (in each 5 mL = 1 teaspoon)	Purposes
Chlorpheniramine maleate, USP 1 mg	Antihistamine
Dextromethorphan HBr, USP 7.5 mg	Cough suppressant

Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
 - runny nose
 - itchy, watery eyes
 - sneezing
 - itching of the nose or throat

Warnings

Do not use

- to sedate a child or to make a child sleepy
- 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

- trouble urinating due to an enlarged prostate gland
- glaucoma
- a cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma, chronic bronchitis or emphysema

Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers.

When using this product

- do not use more than directed
- marked drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

Stop use and ask a doctor if 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 4 doses in any 24-hour period
- mL=mililiter; tsp=teaspoonful

Age (years)	Dose
Under 6	do not use
6-11	2 teaspoons (10 mL) every 6 hours
12 and over	4 teaspoons (20 mL) every 6 hours

Other information

- each teaspoon (5 mL) contains: sodium 1 mg
- store at 15-30°C (59-86°F)
- measure only with dosage cup provided
- do not use if printed bottle wrap is missing or broken.

Inactive ingredients

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

Questions

1-800-858-3889

Manufactured by: DLC Laboratories, Inc. Paramount, CA 90723 USA Long Acting TRUSTED SINCE 1978

Children's

Bactimicina

Milti-Symptoms • Liquid Ages 6+ Years

Cough & Cold

Dextromethorphan HBr (Cough Suppressant)

Chlorpheniramine Maleate (Antihistamine)

Relieves Cough up to 8hrs • Sneezing • Runny Nose

Alcohol Free

Natural Strawberry Flavor

4 FL OZ (118 mL)

Drug Facts

Active ingredients Purposes (in each 5 mL = 1 teaspoon)

Chlorpheniramine maleate, USP 1 mg....Antihistamine Dextromethorphan HBr, USP 7.5 mg.....Cough suppressant

temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: runny nose

■sneezing ■itchy, watery eyes ■itching of the nose or throat

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Long Acting

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Children's Bactimicina

Multi-Symptom • Liquid

Ages **6+** Years

Cough & Cold

Relieves Cough up to 8 hrs • Sneezing • Runny Nose



Drug Facts (continued)

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Ask a doctor before use if you have

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Manufactured by/Fabricado por

Propósitos

@delacruzproducts

f @DLClaboratories

Drug Facts Datos Medicinales

Peel Here evante Aquí

P0173-GYY

Do not use if printed bottle wrap is missing or broken.

drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping

When using this product

do not use more than directed marked drowsiness may occur ■avoid alcoholic drinks ■alcohol, sedatives and tranquilizers may increase drowsiness

be careful Stop use and ask a doctor if cough lasts more than when driving a motor vehicle or operating machinery

excitability may occur, especially in children

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 days, comes back, or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition.

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■each teaspoon (5 mL) contains: sodium 1 mg ■store at 5-30°C (59-86°F) ■measure only with dosage cup provided

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

Datos Medicinales

.Antihistamínico Naleato de clorfenamina, USP 1 ma. Dextrometorfano HBr, USP 7.5 ma

'en cada 5 mL = 1 cucharadita

naredientes activos

ocurrir por un restrio ■ alivia temporalmente estos sintomas de rinitis alérgica o algunas alergias de las vías Usos = alivia temporalmente la tos causada por una rritación leve de garganta y bronquios, como puede

■picazón de ojos, ojos llorosos ■estornudos respiratorias superiores:

goteo nasal comezón nasal o de garganta

Advertencias No usar

para sedar a un niño o hacerlo sentir soñoliento

YY2-6YY

Datos Medicinales (continuación)

 si está tomando un medicamento recetado inhibidor de Si no sabe si su receta contiene IMAO, pregúntele para la depresión, afecciones psiquiátricas o emocionales o enfermedad de Parkinson), o en un período de 2 semanas a un médico o farmacéutico antes de usar este producto después de haber suspendido el uso del medicamento monoaminooxidasa (IMAO) (ciertos medicamentos

Drug Facts (continued)

glándula prostática e glaucoma e tos que ocurre con demasiada flema (mucosidad) e problemas respiratorios o tos crónica que perdura o se produce por fumar, asma Pregúntele a un médico antes de usar si usted tiene problemas al orinar debido al agrandamiento de la bronquitis crónica o enfisema

Consulte con un médico o farmacéutico antes de

usarlo si usted está tomando sedantes o tranquilizantes Cuando use este producto ■no exceda la dosis

■tenga cuidado si conduce u opera maquinaria ■ puede recomendada puede provocar mucha somnolencia pevite tomar alcohol el alcohol, los sedantes o los tranquilizantes pueden incrementar la somnolencia provocar excitabilidad, especialmente en niños.

Deje de usar y consulte con un médico si

profesional de la salud antes de usar. Manténgase fuera del alcance de los niños. En caso de sobredósis, pida ayuda médica o contáctese con un Centro de Control de tos dura más de 7 días, regresa o está acompañada por fiebre, salpullido o dolor de cabeza persistente. Estos podrían ser síntomas de una afección grave. Si está embarazada o lactando, consulte con un a

Instrucciones

Envenenamientos inmediatamente

■no tomar más de 4 dosis en un período de 24 horas mL = millitro

2 cucharaditas (10 mL) cada 6 horas 4 cucharaditas (20 mL) cada 6 horas no lo use 12 y mayores Menor de 6 Edad 6-11

■cada cucharadita (5 mL) contiene: 1 mg de sodio ■consérvese a 15-30°C (59-86°F) ■use únicamente vaso dosificador incluido Otra información

Ingredientes inactivos ácido cítrico, FD&C rojo no. 40, saborizante, glicerina, propilenglicol, agua purificada, benzoato de sodio, sorbitol, sucralosa

BACTIMICINA CHILDRENS COUGH AND COLD

chlorpheniramine maleate, dextromethorphan hydrobromide liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:24286-1551

Route of Administration ORAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
CHLORPHENIRAMINE MALEATE (UNII: V1Q0090J9Z) (CHLORPHENIRAMINE - UNII: 3U6I01965U)	CHLORPHENIRAMINE MALEATE	1 mg in 5 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	7.5 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)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SORBITOL (UNII: 506T60A25R)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		
GLYCERIN (UNII: PDC6A3C0OX)		

Product Characteristics		
Color	red (Red)	Score
Shape		Size
Flavor	STRAWBERRY (Natural Strawberry flavor)	Imprint Code
Contains		

F	Packaging			
#	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24286- 1551-4	1 in 1 BOX	10/23/2009	
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date

OTC monograph final	part341	10/23/2009

Labeler - DLC Laboratories, Inc. (093351930)

Establishment			
Name	Address	ID/FEI	Business Operations
DLC Laboratories, Inc.		093351930	manufacture(24286-1551) , label(24286-1551)

Revised: 11/2022 DLC Laboratories, Inc.