COUGH RELIEF ADULT- dextromethorphan hbr liquid Dolgencorp, Inc. (DOLLAR GENERAL & REXALL)

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.

Drug Facts

Active ingredient (in each 5 mL)

Dextromethorphan HBr 15 mg

Purpose

Cough suppressant

Uses

- temporarily relieves
 - cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
 - the impulse to cough to help you get to sleep

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

- cough that occurs with too much phlegm (mucus)
- chronic cough that lasts as occurs with smoking, asthma, or emphysema

When using this product,

do not use more than directed

Stop use and ask a doctor if

cough lasts more than 7 days, cough comes back, or occurs by fever, rash or headache that lasts. 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 (1-800-222-1222) right away.

Directions

- take recommended dosage or as directed by a doctor
- measure only with dosing cup provided. Do not use any other dosing device
- keep dosing cup with product
- shake well before using
- mL=milliliter

age	dose	
adults and children 12 years and	10 mL every 6-8 hours not to exceed 4 doses in	
over	24 hours,	
children 6 to 11 years of age	5 mL every 6-8 hours not to exceed 4 doses in 24	
children o to 11 years or age	hours	
children under 6 years of age	do not use	

Other information

- each 5 mL contains: sodium 3 mg
- store between 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

anhydrous citric acid, disodium EDTA, FD&C yellow #6, flavor, glycerin, high fructose corn syrup, hydroxyethyl cellulose, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sucralose, sucrose, xanthan gum

Questilons or comments?

Call 1-888-309-9030

Principal Display Panel

For Ages 6 Years and over

Adult

Cough Relief

Dextromethorphan HBr 15 mg

Cough suppressant

Relieves:

- Cough
- Minor throat and bronchial irritation
- impulse to cough

6-8 Hour Relief

Alcohol free

FL OZ (mL)

Orange flavor

Dosing Cup Included

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF PRINTED SAFETY SEAL AROUND BOTTLE OR UNDER CAP IS BROKEN OR MISSING.

DISTRIBUTED BY DOLGENCORP, LLC 100 MISSION RIDGE GOODLETTSVILLE, TN 37072

Package Label



COUGH RELIEF ADULT

dextromethorphan hbr liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:55910-309

Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) DEXTROMETHORPHAN - UNII:7355X3ROTS) DEXTROMETHORPHAN | 15 mg | in 5 mL

Inactive Ingredients	
Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
WATER (UNII: 059QF0KO0R)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GLYCERIN (UNII: PDC6A3C0OX)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
HYDROXYETHYL CELLULOSE (1500 MPA.S AT 1%) (UNII: L605B5892V)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SUCROSE (UNII: C151H8M554)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics				
Color		Score		
Shape		Size		
Flavor	ORANGE	Imprint Code		
Contains				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:55910- 309-04	1 in 1 BOX	08/31/2015		
1		118 mL in 1 BOTTLE, PLASTIC; 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	08/31/2015		

Labeler - Dolgencorp, Inc. (DOLLAR GENERAL & REXALL) (068331990)

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