COUGH RELIEF COUGH SUPPRESSANT- dextromethorphan hbr liquid
QUALITY CHOICE (Chain Drug Marketing Association)

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

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

Stop use and ask a doctor if

cough lasts more than 7 days, comes back, or occurs with fever, rash or headache that lasts. These could be signs of a serious condition.

When using this product

- **do not use more than directed**

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 over	10 mL every 6-8 hours no to exceed 4 doses in 24 hours
children 6 to 11 years of age	5 mL every 6-8 hours not to exceed 4 doses in 24 hours
children under 6 years of age	do not use

Other information

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

Inactive ingredients

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

Questions or comments?

Call **1-248-449-9300 Monday-Friday 9AM-5PM EST**

Principal Display Panel

COUGH RELIEF

Cough Suppressant

Dextromehtorphan HBr 15 mg

Cough Suppressant

For Ages 6 & Over

6-8 Hour Relief

Alcohol Free

Orange Flavor

FL OZ (mL)

Contains no fever reducer or pain reliever

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 C.D.M.A., Inc,©

43157 W. Nine Mile

Novi, MI 48376-0995

www.qualitychoice.com

Package Label

QUALITY CHOICE Cough Relief

COUGH RELIEF COUGH SUPPRESSANT

dextromethorphan hbr liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-957

Route of Administration	ORAL

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SUCROSE (UNII: C151H8M554)	
XANTHAN GUM (UNII: TTV12P4NEE)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYDROXYETHYL CELLULOSE (1500 MPA.S AT 1%) (UNII: L605B5892V)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-957-04	1 in 1 BOX	07/31/2015	
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 Drug	M012	07/31/2015	

Labeler - QUALITY CHOICE (Chain Drug Marketing Association) (011920774)

Registrant - P & L Development, LLC (079765031)

Revised: 4/2024 QUALITY CHOICE (Chain Drug Marketing Association)