LONG ACTING COUGH SOFTGELS- dextromethorphan hydrobromide capsule, liquid filled

Chain Drug Marketing Association, Inc.

QC® QUALITY CHOICE Adult Long Acting Cough Softgels

Drug Facts

Active ingredient (in each softgel)

Dextromethorphan HBr, USP 15 mg

Purpose

Cough Suppressant

Use

temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold.

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

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

Stop use and ask a doctor if cough lasts for more than 7 days, comes back, or is accompanied by fever, rash, or persistent headache. These can be signs of a serious condition.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. Overdose warning: In case of overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as children even if you do not notice any signs or symptoms.

Directions

do not take more than 8 softgels in any 24-hour period this adult product is not intended for use in children under 12 years of age

adults and	take 2 softgels
children	every
12 years and	6 to 8 hours, as
older	needed
children under	do not use
12 years	

Other information

- store at 20°-25°C(68°-77°F)
- avoid excessive heat above 40°C(104°F)
- protect from light

Inactive ingredients

edible white ink, FD&C blue #1, FD&C red #40, gelatin, glycerin, isopropyl alcohol, medium chain triglycerides, polyethylene glycol, povidone, propyl gallate, propylene glycol, purified water, sorbitol sorbitan solution

Questions or comments?

1-888-577-8033 Monday - Friday 8am - 4pm EST

Compare to the active ingredient in Robitussin® CoughGels®*

Relieves cough for up to 8 Hours Non-Drowsy

READ AND KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SEAL UNDER CAP IS BROKEN OR MISSING FROM BOTTLE

Distributed by: C.D.M.A., Inc.© 43157 W 9 Mile Rd Novi, MI 48375 www.qualitychoice.com Product of UAE

Packaged and Quality Assured in the USA

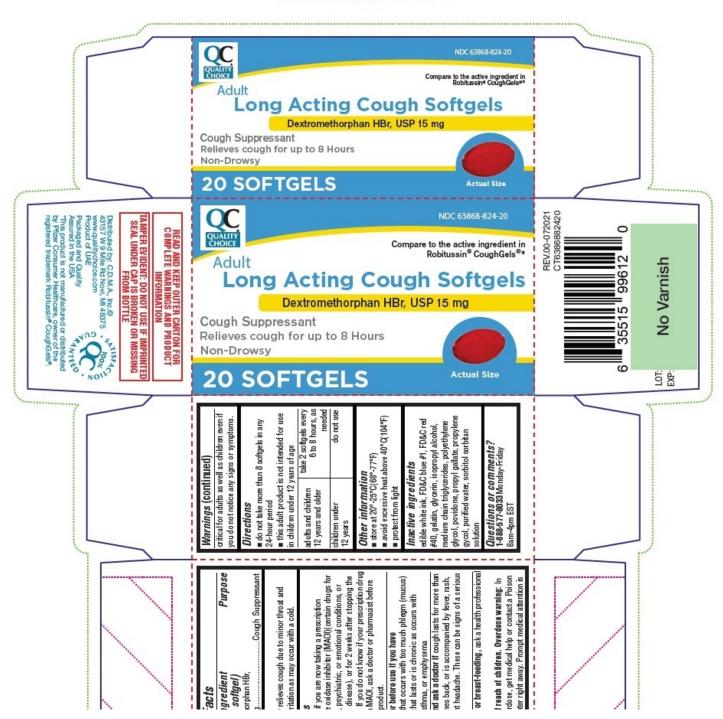
*This product is not manufactured or distributed by Pfizer Consumer Healthcare, owner of the registered trademark Robitussin® CoughGels®.

100% QC SATISFACTION GUARANTEED

Packaging



OUTER PACKAGE LABEL



DRUG FACTS TABLE

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Warnings (continued)

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Directions

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adults and children 12 years and older	take 2 softgels every 6 to 8 hours, as needed
children under	do not use
12 years	

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LONG ACTING COUGH SOFTGELS

dextromethorphan hydrobromide capsule, liquid filled

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:63868-824

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
	DEXTROMETHORPHAN HYDROBROMIDE	15 mg

Inactive Ingredients		
Ingredient Name	Strength	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)		
GLYCERIN (UNII: PDC6A3C0OX)		
ISOPROPYL ALCOHOL (UNII: ND2M416302)		
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)		
PROPYL GALLATE (UNII: 8D4SNN7V92)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SORBITOL (UNII: 506T60A25R)		
SORBITAN (UNII: 6O92ICV9RU)		

Product Characteristics			
Color	red	Score	no score
Shape	OVAL (oblong)	Size	14mm
Flavor		Imprint Code	778
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-824- 20	1 in 1 CARTON	05/03/2022	
1		20 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	05/03/2022	

Labeler - Chain Drug Marketing Association, Inc. (011920774)