COUGH RELIEF- dextromethorphan hbr capsule, liquid filled WALGREEN COMPANY

Walgreens Non-Drowsy Wal-Tussin Cough Liquid Gels

Drug Facts

Active ingredient (in each softgel)

Dextromethorphan HBr, USP 15mg

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 doctor if cough lasts for 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. (1-800-222-1222)

Directions

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

12 years and over	every o to o nours,		
12 years and over	as needed		
children under 12	do not use		
years	do not use		

Other information

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

Inactive ingredients

FD&C Red No. 40, gelatin, glycerin, polyethylene glycol 400, propylene glycol, povidone, sorbitol sorbitan solution, titanium dioxide

Questions or comments?

1-888-333-9792

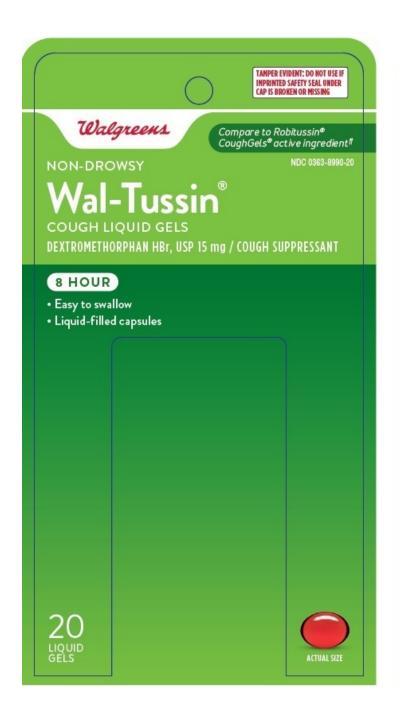
PRINCIPAL DISPLAY PANEL

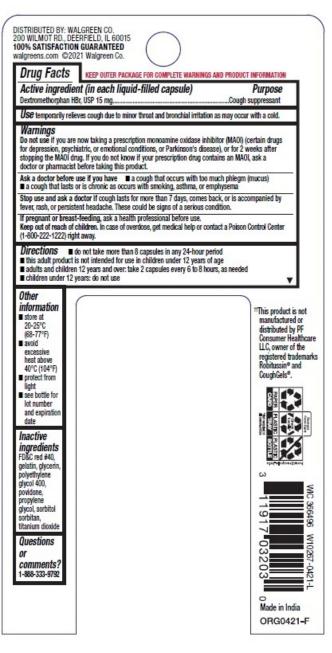
Compare to Robitussin® Cough Gels® active ingredient*

Cough Relief

Dextromethorphan HBr 15mg

20 softgels





COUGH RELIEF

dextromethorphan hbr capsule, liquid filled

D	ro	dı	ıct	Inf	rm	ati	on	
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:0363-8990

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH)	DEXTROMETHORPHAN	15 mg

Inactive Ingredients					
Ingredient Name	Strength				
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)					
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)					
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)					
WATER (UNII: 059QF0KO0R)					
GELATIN (UNII: 2G86QN327L)					
GLYCERIN (UNII: PDC6A3C0OX)					
SORBITOL (UNII: 506T60A25R)					
FD&C RED NO. 40 (UNII: WZB9127XOA)					
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)					

Product Characteristics					
Color	red	Score	no score		
Shape	OVAL	Size	13mm		
Flavor		Imprint Code	404;904		
Contains					

F	Packaging								
#	tem Code	Package Description	Marketing Start Date	Marketing End Date					
1	NDC:0363- 8990-20	1 in 1 PACKAGE	12/22/2020						
1		20 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	12/22/2020				

Labeler - WALGREEN COMPANY (008965063)

Revised: 12/2023 WALGREEN COMPANY