

**ROBITUSSIN MAXIMUM STRENGTH COUGH PLUS CHEST CONGESTION DM-
dextromethorphan hydrobromide and guaifenesin capsule, liquid filled
Haleon US Holdings LLC**

Drug Facts

Active ingredients (in each liquid-filled capsule)

Dextromethorphan HBr, USP 10 mg

Guaifenesin, USP 200 mg

Purposes

Cough suppressant

Expectorant

Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes

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)
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis, or emphysema

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

age	dose
adults and children 12 years and over	2 capsules every 4 hours
children under 12 years	do not use

Other information

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

Inactive ingredients

FD&C Red No. 40, gelatin, glycerin, light mineral oil, pharmaceutical ink, polyethylene glycol, povidone, propylene glycol, purified water, sorbitol sorbitan solution

Questions or comments?

call weekdays from 9 AM to 5 PM EST at **1-800-762-4675**

Made in Canada

For most recent product information, visit www.robitussin.com

Distributed by: Pfizer, Madison, NJ 07940 USA

PRINCIPAL DISPLAY PANEL

ADULT

Robitussin®

**MAXIMUM
STRENGTH**

**Cough+Chest
Congestion DM**

**DEXTROMETHORPHAN HBr (Cough Suppressant)
GUAIFENESIN (Expectorant)**

**DM
MAX**

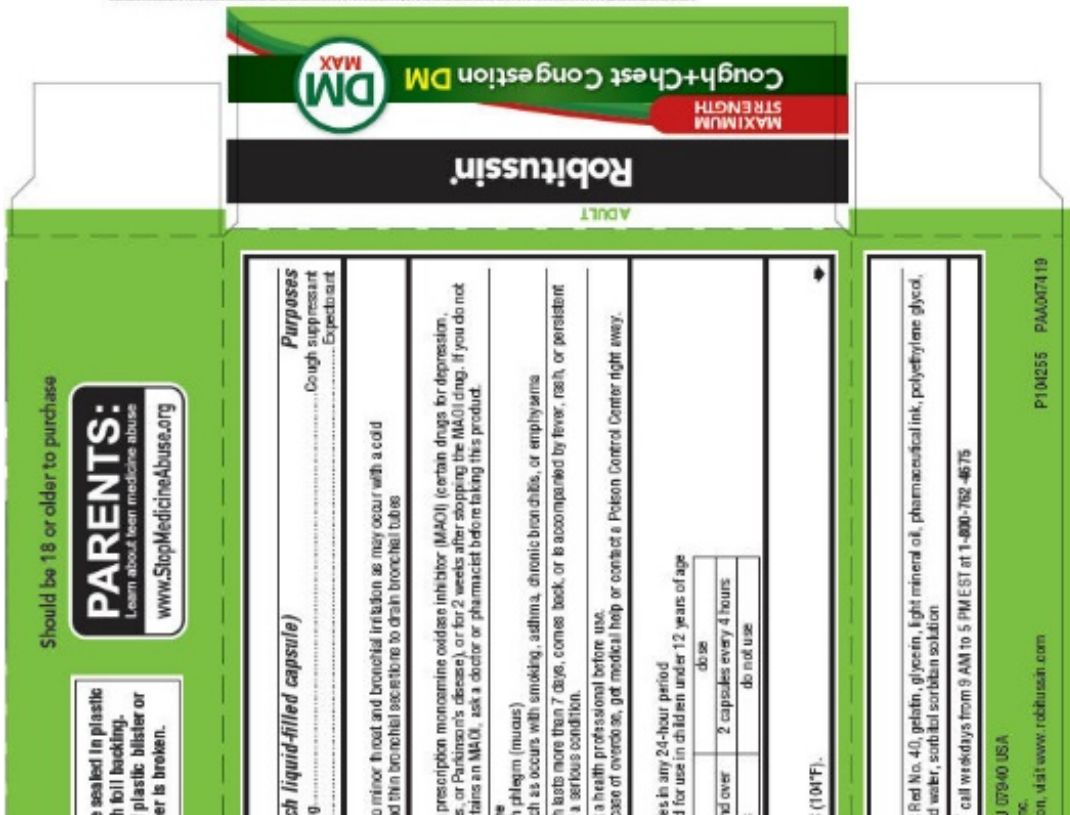
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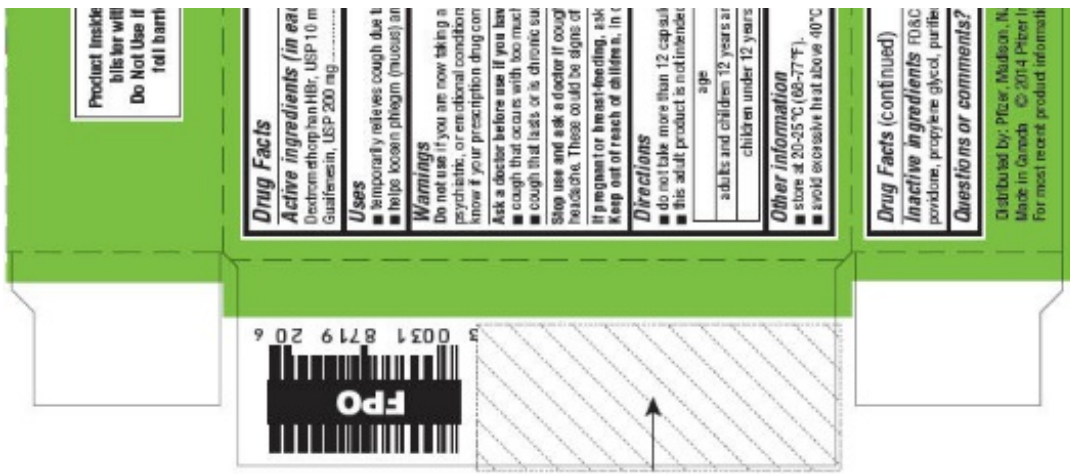
- Cough
- Mucus

Non-Drowsy

For Ages 12 & Over

20
LIQUID-FILLED
CAPSULES





ROBITUSSIN MAXIMUM STRENGTH COUGH PLUS CHEST CONGESTION DM

dextromethorphan hydrobromide and guaifenesin capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0031-8719
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
SORBITAN (UNII: 6O92ICV9RU)	

Product Characteristics

Color	RED (clear red)	Score	no score
Shape	OVAL (oblong)	Size	16mm
Flavor		Imprint Code	R

Contains**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0031-8719-10	5 in 1 CARTON	05/02/2016	
1		2 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:0031-8719-20	10 in 1 CARTON	05/23/2013	
2		2 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:0031-8719-31	15 in 1 CARTON	05/23/2013	12/31/2018
3		2 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:0031-8719-70	500 in 1 BOX	05/23/2013	12/31/2018
4		2 in 1 POUCH; 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	05/23/2013	

Labeler - Haleon US Holdings LLC (079944263)

Revised: 3/2024

Haleon US Holdings LLC