

**ROBITUSSIN LONG-ACTING COUGH SOFT CHEWS- dextromethorphan hydrobromide tablet, chewable**  
**Haleon US Holdings LLC**

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***Drug Facts***

***Active ingredient (in each chewable tablet)***

Dextromethorphan equivalent to dextromethorphan HBr 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**

- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, 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 4 doses (8 chewable tablets) 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	take 2 chewable tablets every 6 to 8 hours, as needed
children under 12 years	do not use

***Other information***

- each chewable tablet contains:** potassium 10 mg, sodium 6 mg
- store at a controlled room temperature 20-25°C (68-77°F)

### ***Inactive ingredients***

carboxymethylcellulose calcium, carnauba wax, corn syrup solids, crospovidone, FD&C red no. 40, glycerin, mannitol, natural and artificial flavors, pregelatinized starch, simethicone, sodium gluconate, sorbitol, sorbitol solution, sucralose, sucrose, whole dry milk

### ***Questions or comments?***

call weekdays from 8 AM to 6 PM EST at **1-800-245-1040**

### **Additional Information**

**Do Not Use if seal under bottle cap imprinted with  
“SEALED for YOUR PROTECTION” is broken or missing.**

### **PARENTS:**

Learn about teen medicine abuse

**[www.StopMedicineabuse.org](http://www.StopMedicineabuse.org)**

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### **PRINCIPAL DISPLAY PANEL**

#### **NEW! ADULT HALEON**

**Robitussin**

**Long-Acting Cough**

**Soft CHEWS**

**DEXTROMETHORPHAN HBr 15 mg  
(Cough Suppressant)**

**FAST relief ANYWHERE**

**UP TO 8HRCOUGH RELIEF**

Chew tablets completely before swallowing

Berry flavor

**10 CHEWABLE TABLETS**

208802 Front Label



ROBITUSSIN LONG-ACTING COUGH SOFT CHEWS			
dextromethorphan hydrobromide tablet, chewable			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0031-9310
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
Inactive Ingredients			
Ingredient Name			Strength
CARBOXYMETHYLCELLULOSE CALCIUM (UNII: UTY7PDF93L)			
CARNAUBA WAX (UNII: R12CBM0EIZ)			
CORN SYRUP (UNII: 9G5L16BK6N)			
CROSPVIDONE, UNSPECIFIED (UNII: 2S7830E561)			
FD&C RED NO. 40 (UNII: WZB9127XOA)			
GLYCERIN (UNII: PDC6A3C0OX)			
MANNITOL (UNII: 3OWL53L36A)			
STARCH, CORN (UNII: O8232NY3SJ)			

<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)				
<b>SODIUM GLUCONATE</b> (UNII: R6Q3791S76)				
<b>SORBITOL</b> (UNII: 506T60A25R)				
<b>SORBITOL SOLUTION</b> (UNII: 8KW3E207O2)				
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)				
<b>SUCROSE</b> (UNII: C151H8M554)				
<b>COW MILK</b> (UNII: 917J3173FT)				
<b>Product Characteristics</b>				
<b>Color</b>	pink (darker pink speckles)		<b>Score</b>	no score
<b>Shape</b>	ROUND		<b>Size</b>	21mm
<b>Flavor</b>	BERRY		<b>Imprint Code</b>	R
<b>Contains</b>				
<b>Packaging</b>				
<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:0031-9310-10	10 in 1 BOTTLE; Type 0: Not a Combination Product	06/19/2024	
2	NDC:0031-9310-20	20 in 1 BOTTLE; Type 0: Not a Combination Product	06/19/2024	
<b>Marketing Information</b>				
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>		<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M012		06/19/2024	

**Labeler** - Haleon US Holdings LLC (079944263)