ROBOCOUGH- dextromethorphan hbr liquid DXM Pharmacuetical, Inc.

Active ingredient

(in each 3 mL) (one dose) Dextromethrophan HBr, 30 mg

Purpose

Cough suppressant

Uses

• Temporarily relieves cough due to minor throat and bronchial irritation as may occur with the common cold.

Warning

Do not useif 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 is accompanied by excessive phlegm (mucus), or
- a persistent or chronic cough such as occurs with smoking, asthma, or emphysema.

Stop use and ask a doctor ifa cough persists for more than 1 week, tends to recur, or is accompanied by fever, rash, or persistent headache. A persistent cough may be a sign 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 exceed 120 milligrams (4 doses) in any 24-hour period
- measure only 3 mL with dosing syringe provided
- only use this product according to these directions

age dose

adults and children 12 years and over

3mL every 6 to 8 hours

children under 12 years

Other information

- **TAMPER EVIDENT**: Do not use if seal on the bottle is broken.
- store at 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

Citric acid, glycerin, grape flavor, propylene glycol, purified water, sodium benzoate, sodium citrate, stevia, sucralose, sucrose

Questions or comments?

833-289-7626 between 10 a.m. to 4 p.m. CST, Monday-Friday

Product label



ROBOCOUGH

dextromethorphan hbr liquid

Product Information							
Product Type	HUMAN OTC DRUG	Item Code (Source	e)	NDC:7159	98-102		
Route of Administration	ORAL						
Active Ingredient/Active	Maiatur						
Active Ingredient/Active Moiety							
Ingre	Ba	sis of Str	rength	Strength			

Inactive Ingree	dients					
3	Strength					
ANHYDROUS CITRI	C ACID (UNII:)	Ingredient Name (F417D3PSL)			-	
GLYCERIN (UNII: PD	C6A3C0OX)					
PROPYLENE GLYCO	L (UNII: 6DC9	Q167V3)				
NATER (UNII: 059QF	OKOOR)					
ODIUM BENZOAT	E (UNII: 0J245F	E5EU)				
SODIUM CITRATE (UNII: 1Q73Q2JI	JLR)				
STEVIA LEAF (UNII:	6TC6NN0876)					
SUCRALOSE (UNII: 9	96K6UQ3ZD4)					
SUCROSE (UNII: C15	51H8M554)					
Product Chara	cteristics					
Color			Score			
Shape			Size			
lavor GRAPE		Imprint Code				
Contains						
Packaging						
# Item Code	Pa	ckage Description		Marketing Start Date	Marketing End Date	
1 NDC:71598-102- 40	L in 1 CARTON			10/01/2023		
	120 mL in 1 BOTTLE; Type 0: Not a Combination Product					
Markating	nformat	ion				
Marketing I						
Marketing	Applica	tion Number or Mo	onograph	Marketing Start	Marketing End	
Category		Citation		Date	Date	

Labeler - DXM Pharmacuetical, Inc. (080748277)

Registrant - DXM Pharmacuetical, Inc. (080748277)

Revised: 1/2025

DXM Pharmacuetical, Inc.