

ROBO COUGH- dextromethorphan hbr liquid
DXM Pharmacuetical, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Robo Cough

In each 3ml (one dose)

Dextromethrophan HBr 30 mg.....cough suppressant

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

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 is accompanied by excessive phlegm (mucus), or
- a persisten or chronic cough such as occurs with smoking, asthma, or emphysema.

Stop use and ask a doctor if a 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

- only use this product according to these directions or as directed by a doctor.
- do not exceed 120 milligrams in any 24-hour period
- measure only 3mL with dosing cup provided
- keep dosing cup with product
- mL= milliliter
- this adult product is not intended for use in children under 12 years of age.

age	dose
adults and children 12 years and over	3mL every 6 to 8 hours
children under 12 years	do not use

Other information

- store at 20-25* (68-77°F). Do not refrigerate.
- **TAMPER EVIDENT:** Do not use if printed shrink band on bottle is broken or missing.

Inactive ingredients high fructose corn syrup, methyl anthranilate (artificial grape flavor), phosphoric acid, propylene glycol, purified water, sodium benzoate (preservative), sucralose

Questions or comments?

Call 1-833-289-7626 between 10 a.m. to 4 p.m. CST, Monday through Friday or visit our website www.RoboCough.com to report serious adverse events associated with the use of this product. Please

call a doctor for medical advice.
 Dextromethorphan HBr.....30mg
 Distributed by:
 DXM Pharmaceutical, Inc.
 2717 Commercial Center Blvd.,
 Suite E200
 Katy, TX 77494
 www.RoboCough.com
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GRAPE MAXIMUM COUGH RELIEF 15 DOSES

GRAPE CONCENTRATED COUGH RELIEF 15 DOSES

GRAPE MAXIMUM COUGH RELIEF 15 DOSES

GRAPE CONCENTRATED COUGH RELIEF 15 DOSES

ADULT Robo Cough DO NOT USE IF SEAL IS BROKEN

ADULT Robo Cough DO NOT USE IF SEAL IS BROKEN

up to 8 HOUR COUGH RELIEF per dose

15 Doses per bottle 30 mg per dose of Dextromethorphan HBr

Distributed by DXM Pharmaceutical, Inc.,
 2717 Commercial Center Blvd.,
 Suite E200, Katy, TX 77494
www.RoboCough.com

POWERFUL Robo Cough SUPPRESSANT

15 Doses per bottle

450mg Dextromethorphan HBr

MADE IN USA

SEVERE COUGH Drug Facts

Active ingredient Purpose (in each 3 mL) (one dose)
 Dextromethorphan HBr,
 30 mg.....Cough suppressant

Uses ■ temporarily relieves cough due to minor throat and bronchial irritation as may occur with the common cold.

Warnings Do not use if you are now taking a prescription

Drug Facts (continued)
PEEL and LIFT Label!

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NOT RECOMMENDED FOR HOUSEHOLDS WITH CHILDREN. PATENT PENDING 1.52 fl oz (45 mL) NOT RECOMMENDED FOR HOUSEHOLDS WITH CHILDREN.

ROBO COUGH

dextromethorphan hbr liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71598-000
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg in 3 mL
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Inactive Ingredients

Ingredient Name	Strength
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
METHYL ANTHRANILATE (UNII: 981I0C1E5W)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0K00R)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	GRAPE	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71598-000-00	45 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/19/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	03/19/2018	

Labeler - DXM Pharmaceutical, Inc. (080748277)

Registrant - DXM Pharmaceutical, Inc. (080748277)

Establishment

Name	Address	ID/FEI	Business Operations
Woodfield Pharmaceutical, LLC		079398730	manufacture(71598-000)

Establishment

Name	Address	ID/FEI	Business Operations
DXM Pharmaceutical, Inc.		080748277	label(71598-000)