

ADULT COUGH RELIEF DM- dextromethorphan hbr liquid

Safeway, 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.

Drug Facts

Active ingredient (in each 10 mL)

Dextromethorphan HBr 30 mg

Purpose

Cough suppressant

Uses

- 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 (1-800-222-1222) right away.

Directions

- do not take more than 4 doses in any 24-hour period
- measure only with dosing cup provided. Do not use any other dosing device
- keep dosing cup with product
- mL = milliliter
- this adult product is not intended for use in children under 12 years of age
- adults and children 12 years and over 10 mL every 6 to 8 hours
- children under 12 years of age: do not use

Other information

- store between 20-25°C (68°-77°F). Do not refrigerate.

Inactive ingredients

anhydrous citric acid, FD&C red #40, dextrose, ethyl alcohol, flavor, glycerin, high fructose corn syrup, menthol, purified water, saccharin sodium, sodium benzoate

Questions or comments?

Call 1-888-723-3929 Monday-Friday 7AM-6PM PST

Principal Display Panel**Adult Cough Relief DM**

Dextromethorphan HBr 30 mg-Cough Suppressant

ORIGINAL FLAVOR

Compare to Robitussin® Lingerin Cold Long-Acting Cough active ingredient*

- For ages 12 & over
- Alcohol 1.4%
- Dosing cup included
- Non-Drowsy

FL OZ (mL)

*This product is not manufactured or distributed by Pfizer Consumer Healthcare, distributors of Robitussin® Lingerin Cold Long-Acting Cough.

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF PRINTED SAFETY SEAL AROUND BOTTLE OR UNDER CAP IS BROKEN OR MISSING.

DISTRIBUTED BY

BETTER LIVING BRANDS LLC

P.O BOX 99, PLEASANTON, CA 94566-0009

1888-723-392

www.betterlivingbrandsLLC.com

Package Label

Drug Facts (continued)**Questions or comments?**

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OUR PROMISE

QUALITY & SATISFACTION
**100%
GUARANTEED**
OR YOUR MONEY BACK.

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PARENTS:

Learn about teen medicine abuse

www.StopMedicineAbuse.org



Dosing Cup Included

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Long-Acting
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4 FL OZ (118 mL)

NDC 21130-383-04



Quality Guaranteed

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Cough Relief DM**

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S3398

RD 17047



PLD-0323C FC004104

Lot No.:

Exp. Date:

SIGNATURE CARE Adult Cough Relief DM**ADULT COUGH RELIEF DM**

dextromethorphan hbr liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-383
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg in 10 mL

Inactive Ingredients

Ingredient Name			Strength	
ALCOHOL (UNII: 3K9958V90M)				
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
GLYCERIN (UNII: PDC6A3C0OX)				
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)				
MENTHOL (UNII: L7T10EIP3A)				
WATER (UNII: 059QF0KO0R)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
GLUCOSE OXIDASE (UNII: 0T8392U5N1)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-383-04	1 in 1 BOX	06/30/2014	
1		118 mL 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 FINAL		part341	06/30/2014	

Labeler - Safeway, Inc. (009137209)

Revised: 10/2019

Safeway, Inc.