HEALTHY ACCENTS TUSSIN DM- dextromethorphan hydrobromide and guaifenes in liquid DZA Brands LLC

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.

DZA Brands, LLC Tussin DM Drug Facts

Active ingredient (in each 5 mL tsp)

Dextromethorphan HBr, USP 10 mg Guaifenesin, USP 100 mg

Purpose

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 for more than 7 days, comes back, 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 take more than 6 doses in any 24-hour period
- use only with enclosed measuring cup. Do not use any other device.
- 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 teaspoons (tsp) every 4 hours	
children under 12 years	do not use	

Other information

- each teaspoon contains: sodium 3 mg
- store at 20°-25°C (68°-77°F)

Inactive ingredients

anhydrous citric acid, FD&C red no. 40, flavor, glycerin, high fructose corn syrup, menthol, propylene glycol, purified water, sodium benzoate, sodium citrate, sucralose

Questions or comments?

1-866-322-2439

Principal Display Panel

adult tussin DM

cough & chest congestion

cough suppressant (dextromethorphan HBr)

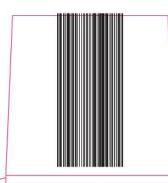
expectorant (guaifenesin)

non-drowsy

for ages 12 & over

Compare to Robitussin® Cough + Chest Congestion DM active ingredients

alcohol free





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age

adults and children

12 years and over

children under 12 years

Other information

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Please include UPC number

and code from package

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DO NOT USE IF PRINTED NECKBAND IS BROKEN OR MISSING

Dosage Cup Included

PARENTS:

Learn about teen medicine abuse

www.StopMedicineAbuse.org

glycol, purified water, sodium benzoate.

dose

2 teaspoons (tsp)

every 4 hours

do not use

Drug Facts (continued)

Directions







NDC 55316-359-34

adult tussin

cough & chest congestion

cough suppressant (dextromethorphan HBr) expectorant (guaifenesin)

> non-drowsy for ages 12 & over

Compare to Robitussin® Cough + Chest Congestion DM

8 FL OZ (237ml)



alcohol free

healthy ** accents**

adult tussin

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LOT NO.

EXP.

: 35934 HA CL



HEALTHY ACCENTS TUSSIN DM

dextromethorphan hydrobromide, guaifenesin liquid

Product Information Product Type

Route of Administration

HUMAN OTC DRUG LABEL

ORAL

Item Code (Source)

DEA Schedule

NDC:55316-359

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
DEXTROMETHORPHAN HYDROBROMIDE (DEXTROMETHORPHAN)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 5 mL		
GUAIFENESIN (GUAIFENESIN)	GUAIFENESIN	100 mg in 5 mL		

Inactive Ingredients		
Ingredient Name	Strength	
ANHYDRO US CITRIC ACID		
FD&C RED NO. 40		
GLYCERIN		
HIGH FRUCTOSE CORN SYRUP		
PROPYLENE GLYCOL		
WATER		
SODIUM CITRATE		
SO DIUM BENZO ATE		
MENTHO L		
SUCRALOSE		

Product Characteristics			
Color	RED (Orange-Red)	Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55316-359-26	1 in 1 CARTON		
1		118 mL in 1 BOTTLE		
2	NDC:55316-359-34	1 in 1 CARTON		
2		237 mL in 1 BOTTLE		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	01/22/2008	

Labeler - DZA Brands LLC (090322194)

Revised: 11/2012 DZA Brands LLC