

SELECTHEALTH TUSSIN DM- guaifenesin liquid
Axcentria Pharmaceuticals, 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.

Tussin
DM

Drug Facts

Active ingredient (in each 5 mL tsp)	Purposes
Dextromethorphan HBr, USP 10 mg	Cough suppressant
Guaifenesin, USP 100 mg	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 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 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 6 doses in any 24-hour period
- this adult product is not intended for use in children under 12 years of age
- **adults and children 12 years of age and over**
2 teaspoons every 4 hours
- **children under 12 years** do not use

Other information

- each teaspoon contains: **sodium 7 mg**
- store 20 to 25°C (68 to 77°F)

- dosage cup provided

Inactive Ingredients

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

Questions?

Adverse drug event call (888) 933-3222

Distributed by:

Axcentria Pharmaceuticals, LLC.
306 Keystone Drive
Telford, PA 18969 USA

PRINCIPAL DISPLAY PANEL - 240 ML Bottle Carton

***COMPARE TO** the active ingredient
in **ROBITUSSIN PEAK COLD**

Tussin
DM

HealthSelect[®]

Cough & Chest
Congestion DM

Dextromethorphan HBr, USP
Cough Suppressant

Antitussiv-Dextrometorfano HBr
Expectorant

Relieves:

- Cough
- Chest Congestion

NON-DROWSY
AGES 12+

NET 8 FL OZ
(240 ML)
NDC 49743-3001-1

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Axcantia Pharmaceuticals, LLC. Re-Order No. #####
306 Keystone Drive #-###
Telford, PA 18969 USA Rev. #/##

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*This product is not manufactured or distributed by Pfizer owner of the registered trademark Robitussin®



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guaifenesin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49743-3001
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Guaifenesin (UNII: 495W7451VQ) (Guaifenesin - UNII:495W7451VQ)	Guaifenesin	100 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
Menthol (UNII: L7T10EIP3A)	
Citric Acid Monohydrate (UNII: 2968PHW8QP)	
Sodium Benzoate (UNII: OJ245FE5EU)	
Saccharin Sodium (UNII: SB8ZUX40TY)	
Glycerin (UNII: PDC6A3C0OX)	

Product Characteristics

Color	RED	Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49743-3001-1	118 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part341	09/01/2011	

Labeler - Axcentria Pharmaceuticals, LLC. (961871501)**Establishment**

Name	Address	ID/FEI	Business Operations
Axcentria Pharmaceuticals, LLC.		961871501	MANUFACTURE

Revised: 8/2011

Axcentria Pharmaceuticals, LLC.