CURATUSSIN DM- dextromethorphan hbr/guaifenesin liquid Wittman Pharma, Inc.

CuraTUSSIN DM

Directions:

- follow dosage below or use as directed by a physician
- do not take more than 6 doses in any 24-hour period

Age	Dose
adults and children 12 years and over	2 teaspoonfuls every 4 hours
children 6 years to under 12 years	1 teaspoonfuls every 4 hours
children 2 years to under 6 years	½ teaspoonfuls every 4 hours
under 2 years	ask a doctor

Warnings: Do not use if you are now taking a prescription monoamine oxidase inhibitor (MADI)

(certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2

weeks after stopping the MADI drug. If you do not know if your prescription drug contains an MAOI; ask a doctor or pharmacist before taking this product.t

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 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.
- ■you are hypersensitive to any of the ingredients.

If pregnant or breastfeeding, 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.

Inactive Ingredients: Citric Acid, Flavor, Methylparaben, Monoammonium Glycyrrhizinate, Potassium Citrate, Propylene Glycol, Propylparaben, Purified Water, Sorbitol, Sucralose

Uses:

■ helps loosen phlegm(mucus) and thin bronchial secretions to make coughs more productive

■ temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Purpose:

Cough Suppressant and Expectorant

Active Ingredient (in each 5 mL tsp)

Dextromethorphan HBr 10mg

Guaifenesin 100mg



CURATUSSIN DM

dextromethorphan hbr/quaifenesin liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83335-102
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
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)		
AMMONIUM GLYCYRRHIZATE (UNII: 3VRD35U26C)		
PROPYLPARABEN (UNII: Z8IX2SC10H)		
POTASSIUM CITRATE (UNII: EE90ONI6FF)		
METHYLPARABEN (UNII: A218C7H19T)		
WATER (UNII: 059QF0KO0R)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		
SORBITOL (UNII: 506T60A25R)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		

Product Characteristics		
Color		Score
Shape		Size
Flavor	GRAPE	Imprint Code
Contains		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83335- 102-16	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/01/2024	
2	NDC:83335- 102-08	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2024	
3	NDC:83335- 102-04	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/01/2024	

Labeler - Wittman Pharma, Inc. (830980947)

Establishment			
Name	Address	ID/FEI	Business Operations
Wittman Pharma, Inc.		830980947	analysis(83335-102), manufacture(83335-102), label(83335-102)

Revised: 4/2024 Wittman Pharma, Inc.