

WAL TUSSIN ADULT CHEST CONGESTION- guaifenesin solution
Walgreen Company

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.

Walgreen Co. Wal-Tussin® Drug Facts

Active ingredient (in each 10 mL)

Guaifenesin, USP 200 mg

Purpose

Expectorant

Use

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

Warnings

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
- measure only 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	10 - 20 mL every 4 hours
children under 12 years	do not use

Other information

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

Inactive ingredients

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

Questions or comments?

1-800-719-9260

Principal Display Panel

ADULT - NON-DROWSY
 Wal-Tussin®
 CHEST CONGESTION
 GUAIFENESIN 200 mg / EXPECTORANT
 ALCOHOL FREE
 Relieves chest congestion & mucus
 12 years & older
 8 FL OZ (237 mL)
 CHERRY FLAVOR



WAL TUSSIN ADULT CHEST CONGESTION

guaifenesin solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-0310
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)			GUAIFENESIN	200 mg in 10 mL
Inactive Ingredients				
Ingredient Name				Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)				
CAMEL (UNII: T9D99G2B1R)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
GLYCERIN (UNII: PDC6A3C0OX)				
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)				
Product Characteristics				
Color	RED (dark)	Score		
Shape		Size		
Flavor	CHERRY	Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0310-26	1 in 1 CARTON	10/13/1995	02/25/2015
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:0363-0310-34	1 in 1 CARTON	10/30/1995	
2		237 mL in 1 BOTTLE; 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	10/13/1995	

Labeler - Walgreen Company (008965063)