TUSSIN MUCUS AND CHEST CONGESTION ADULT- guaifenesin liquid QUALITY CHOICE (Chain Drug Marketing Association)

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 ingredients (in each 10 mL)

Guaifenesin 200 mg

Purposes

Expectorant

Uses

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 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 6 doses in 24-hours 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 to 20 mL every 4 hours
- children under 12 years: do not use

Other information

- each 10 mL contains: sodium 4 mg
- store between 20-25°C (68°-77°F). Do not refrigerate.

Inactive ingredients

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

Questions or comments?

Call 1-248-449-9300 Monday-Friday 9AM-5PM EST

Principal Display Panel

*Compare to the active ingredient in Robitussin® Mucus + Chest Congestion Original

Adult

Tussin

Mucus + Chest Congestion

Guaifenesin 200 mg Expectorant

Relieves:

Mucus

Chest congestion

For ages 12 & Over

Alcohol Free

Non-Drowsy

Original Formula

FL OZ (mL)

*This product is not manufactured or distributed by Pfizer Consumer Healthcare, distributors of Robitussin® Mucus+Chest Congestion Original.

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

Distributed by: C.D.M.A., Inc.©

43157 W. Nine Mile

Novi, Ml 48376-0995 www.qualitychoice.com

Package Label



QUALITY CHOICE Mucus and Chest Congestion

TUSSIN MUCUS AND CHEST CONGESTION ADULT

guaifenesin liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-859
Route of Administration	ORAL		

	1	ngredient Name	Basis of S	Strength	Strength
GUAIFENESIN (U	JNII: 495W	7451VQ) (GUAIFENESIN - UNII:495W7451V(Q) GUAIFENES IN	1	200 mg in 10 m
nactive Ing	redient	S			
Ingredient Name					Strength
ANHYDROUS CI	TRIC ACI	(UNII: XF417D3PSL)			
GLYCERIN (UNII:	PDC6A3C	0OX)			
NATER (UNII: 05	9QF0KO0I	R)			
SACCHARIN SO	DIUM (UN	I: SB8ZUX40TY)			
SODIUM BENZO	ATE (UNI	: OJ245FE5EU)			
D&C RED NO.					
HIGH FRUCTOS	E CORN S	YRUP (UNII: XY6UN3QB6S)			
CARAMEL (UNII:	T9D99G2I	31R)			
MENTHOL (UNII:	L7T10EIP	3A)			
SUCROSE (UNII:	C151H8M	554)			
SUCROSE (UNII:	C151H8M	554)			
SUCROSE (UNII:	C151H8M	554)			
SUCROSE (UNII: Packaging	C151H8M	554)			
		554) Package Description	Marketing Date		Marketing Enc Date
Packaging		Package Description	-	•	
Packaging # Item Code NDC:63868-	1 in 1 B 237 mL	Package Description	Date	•	Date
Packaging # Item Code 1 NDC:63868- 859-08	1 in 1 B 237 mL	Package Description OX in 1 BOTTLE, PLASTIC; Type 0: Not a ation Product	Date	• 1:	Date
Packaging # Item Code 1 NDC:63868- 859-08 1 NDC:63868- 859-08 1 NDC:63868-	1 in 1 B 237 mL Combin 1 in 1 B 118 mL	Package Description OX in 1 BOTTLE, PLASTIC; Type 0: Not a ation Product	05/31/2015	• 1:	Date 2/31/2024
Packaging # Item Code 1 NDC:63868- 859-08 1 NDC:63868- 859-04	1 in 1 B 237 mL Combin 1 in 1 B 118 mL	Package Description OX in 1 BOTTLE, PLASTIC; Type 0: Not a ation Product OX in 1 BOTTLE, PLASTIC; Type 0: Not a	05/31/2015	• 1:	Date 2/31/2024
Packaging # Item Code MDC:63868- 859-08 1 NDC:63868- 859-04 2 NDC:63868- 859-04 2 NDC:63868- 859-04	1 in 1 B 237 mL Combin 1 in 1 B 118 mL Combin	Package Description OX in 1 BOTTLE, PLASTIC; Type 0: Not a ation Product OX in 1 BOTTLE, PLASTIC; Type 0: Not a ation Product	05/31/2015	• 1:	Date 2/31/2024
Packaging # Item Code 1 NDC:63868- 859-08 1 NDC:63868- 859-04	1 in 1 B 237 mL Combin 1 in 1 B 118 mL Combin	Package Description OX in 1 BOTTLE, PLASTIC; Type 0: Not a ation Product OX in 1 BOTTLE, PLASTIC; Type 0: Not a ation Product	Date 05/31/2015 05/31/2015		Date 2/31/2024

Labeler - QUALITY CHOICE (Chain Drug Marketing Association) (011920774)

Revised: 5/2023

QUALITY CHOICE (Chain Drug Marketing Association)