

TUSSIN EXPECTORANT- guaifenesin liquid
AptaPharma, Inc.

Drug Facts

Active ingredient (in each 5 mL teaspoon)

Guaifenesin, USP 100 mg

Purpose

Expectorant

Uses

- Helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and 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

When using this product

- do not use more than directed

Stop use and ask a doctor if

- cough lasts more than 7 days, comes back, or occurs with 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.

Age	Dose
Adults and children 12 years and over	2-4 teaspoonfuls every 4 hours
Children under 12 years	do not use

Other information

- each 5 mL contains: sodium 3 mg

- store between 20-25 °C (68-77°F)
- do not refrigerate
- dosing cup provided
- **Keep carton for full directions for use**

Inactive ingredients

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

Questions? Call weekdays from 9:30AM to 4:30 PM EST at **1-877-798-5944**

Package Label

AP SAFE

NDC 76281-507-24

***COMPARE TO THE ACTIVE
INGREDIENT IN ROBITUSSIN[®]
MUCUS+CHEST CONGESTION.**

**Adult
Tussin
Expectorant**

**Guaifenesin
Oral Solution, USP
Non-Drowsy**

- **Relieves Chest Congestion**
- **Loosens Mucus**
- **For ages 12 & Over**

4 FL OZ (118 mL)

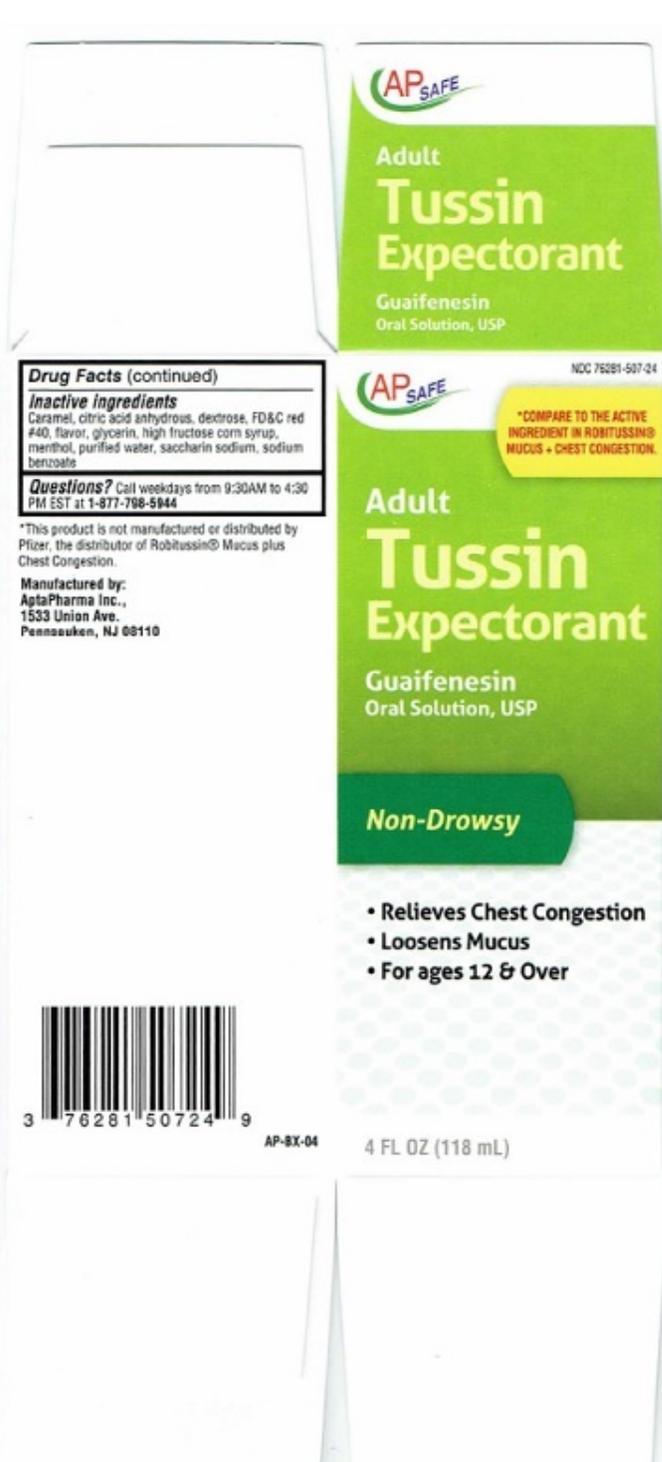
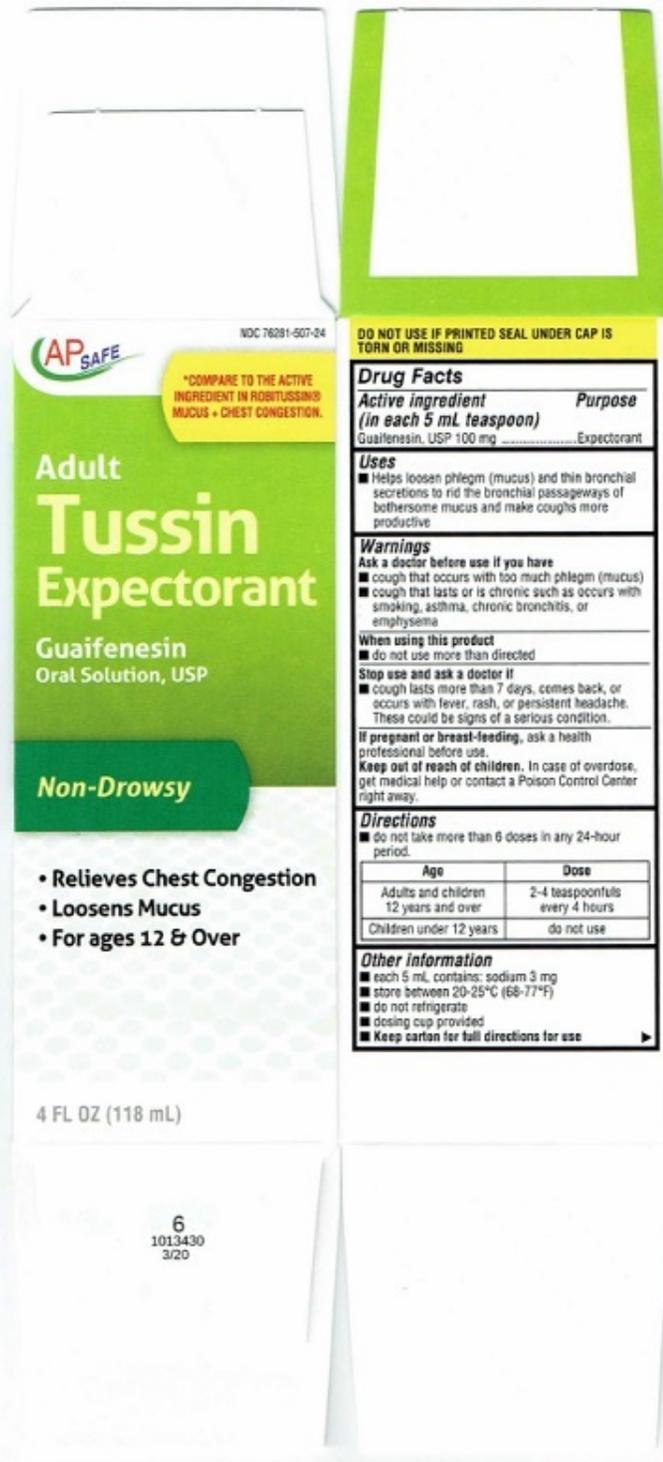
**DO NOT USE IF PRINTED SEAL UNDER CAP IS
TORN OR MISSING**

AP-BX-04

*This product is not manufactured or distributed by Pfizer, the distributor of Robitussin[®] Mucus plus Chest Congestion.

**Manufactured by:
AplaPharma Inc.,
1533 Union Ave.
Pennsauken, NJ 08110**

Carton



DO NOT USE IF PRINTED SEAL UNDER CAP IS TORN OR MISSING

Drug Facts
Active ingredient *Purpose*
 (in each 5 mL teaspoon) Guaifenesin, USP 100 mg Expectorant

Uses
 ■ Helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

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Directions
 ■ do not take more than 6 doses in any 24-hour period.

Age	Dose
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Children under 12 years	do not use

Other information
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 ■ store between 20-25°C (68-77°F)
 ■ do not refrigerate
 ■ dosing cup provided
 ■ Keep carton for full directions for use

Drug Facts (continued)
Inactive ingredients
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AP-BX-04

NDC 76281-507-24



**Adult
Tussin
Expectorant**

Guaifenesin
Oral Solution, USP

Non-Drowsy

- Relieves Chest Congestion
- Loosens Mucus
- For ages 12 & Over

4 FL OZ (118 mL)

DO NOT USE IF PRINTED SEAL UNDER CAP IS BROKEN OR MISSING.

Drug Facts

Active ingredient (in each 5 mL tsp)	Purpose
Guaifenesin, USP 100 mg	Expectorant

Uses ■ Helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

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 1533 Union Ave. Pennsauken, NJ 08110
 AP-LR-04

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EXP:

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TUSSIN EXPECTORANT

guaifenesin liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:76281-507

Route of Administration	ORAL
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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
CARAMEL (UNII: T9D99G2B1R)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
DEXTROSE (UNII: IY9XDZ35W2)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
MENTHOL (UNII: L7T10EIP3A)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

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:76281-507-24	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	09/30/2020	

Labeler - AptaPharma, Inc. (790523323)

Establishment

Name	Address	ID/FEI	Business Operations
AptaPharma, Inc.		790523323	manufacture(76281-507)

