

TUSSIN CF- dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride liquid
AptaPharma Inc.

Drug Facts

Active ingredients
(in each 5 mL tsp)

Dextromethorphan HBr, USP 10 mg
Guaifenesin, USP 100 mg
Phenylephrine HCL, USP 5 mg

Purpose

Dextromethorphan HBr USPCough Suppressant
Guaifenesin USPExpectorant
Phenylephrine HCl, USPNasal Decongestant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes
- temporarily relieves these symptoms occurring with a cold:
 - nasal congestion
 - cough due to minor throat and bronchial irritation.

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 drug contains an MAOI, ask a doctor or pharmacist before taking this product.

Ask a doctor before use if you have

- heart disease ■ high blood pressure
- thyroid disease ■ diabetes
- trouble urinating due to enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema.

Ask a doctor or pharmacist before use if you are taking any other oral nasal decongestant or stimulant.

When using this product do not use more than directed.

Stop use and ask a doctor if

- you get nervous, dizzy or sleepless
- symptoms do not get better within 7 days or are

Multi-Symptom Cold

Dextromethorphan HBr / Guaifenesin /

Phenylephrine HCL

Cough Suppressant / Expectorant /

Nasal Decongestant

Non-Drowsy

Relieves:

- **Cough**
- **Mucus**
- **Nasal Congestion**
- **For Ages 12 and over**

4 FL OZ (118 mL)

CF

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

5

1013430

3/20

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

Manufactured by:

AptaPharma Inc.,

1533 Union Ave,

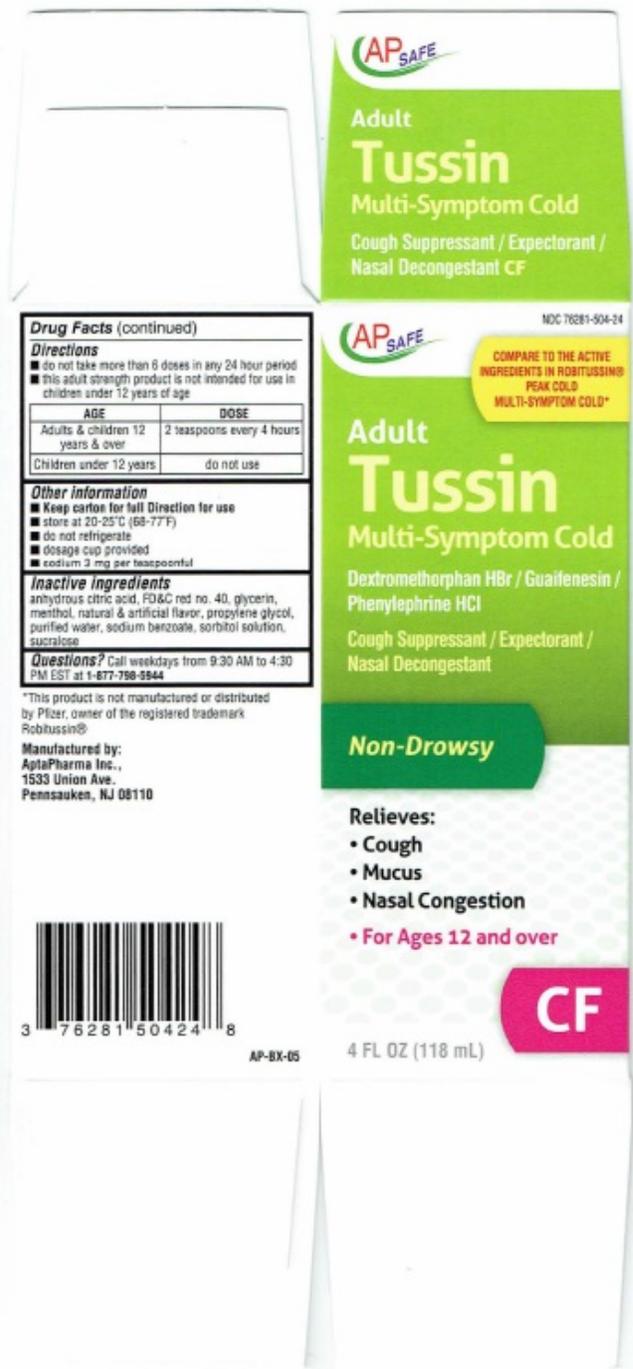
Pennsauken, NJ 08110

Ap·BX·05

Carton



Bottle



AP SAFE
 NDC 76281-504-24
 COMPARE TO THE ACTIVE INGREDIENTS IN ROBITUSSIN® PEAK COLD MULTI-SYMPTOM COLD*

Adult Tussin
 Multi-Symptom Cold
 Dextromethorphan HBr / Guafenesin / Phenylephrine HCl
 Cough Suppressant / Expectorant / Nasal Decongestant

Non-Drowsy

- Relieves:**
- Cough
 - Mucus
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 - For Ages 12 and over

CF

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Ask a doctor or pharmacist before use if you are taking any other oral nasal decongestant or stimulant.

When using this product do not use more than directed.

Stop use and ask a doctor if
 ■ you get nervous, dizzy or sleepless
 ■ symptoms do not get better within 7 days or are accompanied by fever
 ■ 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.

Drug Facts (continued)

Directions
 ■ do not take more than 6 doses in any 24 hour period
 ■ this adult strength product is not intended for use in children under 12 years of age

AGE	DOSE
Adults & children 12 years & over	2 teaspoons every 4 hours
Children under 12 years	do not use

Other information
 ■ **Keep cap on for full Direction for use**
 ■ store at 20-25°C (68-77°F)
 ■ do not refrigerate
 ■ dosage cup provided
 ■ codium 2 mg per teaspoonful

Inactive ingredients
 anhydrous citric acid, FD&C red no. 40, glycerin, menthol, natural & artificial flavor, propylene glycol, purified water, sodium benzoate, sorbitol solution, sucralose

Questions? Call weekdays from 9:30 AM to 4:30 PM EST at 1-877-798-5544

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

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

3 76281 50424 8
 AP-BX-05

AP SAFE
 Adult Tussin
 Multi-Symptom Cold
 Cough Suppressant / Expectorant / Nasal Decongestant CF

AP SAFE
 NDC 76281-504-24
 COMPARE TO THE ACTIVE INGREDIENTS IN ROBITUSSIN® PEAK COLD MULTI-SYMPTOM COLD*

Adult Tussin
 Multi-Symptom Cold
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4 FL OZ (118 mL)



NDC 76281-504-24

Adult

Tussin

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AGE	DOSE
Adults & children 12 years & over	2 teaspoons every 4 hours
Children under 12 years	do not use

Other information ■ store at 20-25°C (68-77°F) ■ do not refrigerate ■ sodium 3 mg per teaspoonful ■ dosage cup provided ■ See carton for full labeling

Inactive ingredients anhydrous citric acid, FD&C red no. 40, glycerin, menthol, natural & artificial flavor, propylene glycol, purified water, sodium benzoate, sorbitol solution, sucralose

Questions? 1-877-798-5944

LOT:

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

AP-LR-05

EXP:

TUSSIN CF

dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76281-504
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 5 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 5 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHOL (UNII: L7T10EIP3A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color	red	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76281-504-24	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/30/2020	

Marketing Information

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

Labeler - AptaPharma Inc. (790523323)

Registrant - AptaPharma Inc. (790523323)

Establishment

Name	Address	ID/FEI	Business Operations
AptaPharma Inc.		790523323	manufacture(76281-504)

Revised: 12/2023

AptaPharma Inc.