TUSSIN CF - dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride liquid
Chain Drug Consortium, LLC

Active ingredients

DRUG FACTS

Active ingredients
(in each 5 mL tsp)
Dextromethorphan HBr, USP 10 mg
Guaifenesin, USP 100 mg
Phenylephrine HCL, USP 5 mg

Purposes

Purposes

Cough Suppressant
Expectorant
Nasal Decongestant

Keep out of reach of children

Keep out of reach of children.

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

Uses

Uses

- help 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

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 Ask a doctor or pharmacist before use

Ask a doctor before use if you have

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an 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

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

If pregnant or breast feeding,

ask a health professional before use.

Directions

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 and children 12 years and over	2 teaspoons every 4 hours
children under 12	do not use

Other information

Other information

- **Keep carton for full Direction for use.**
- store at 20-25 ° C (68-77 ° F)
- do not refrigerate
- dosage cup provided
- sodium 3 mg per teaspoonful

Inactive ingredients

Inactive ingredients

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

Questions?

Questions?

Call weekdays from 9:30 AM to 4:30 PM EST at

1-877-798-5944

Product Label

NDC 68016-126-04

***COMPARE TO THE ACTIVE INGREDIENTS IN ROBITUSSIN® PEAK COLD MULTI-SYMPTOM COLD**

Premier Value ®
Tussin CF

Dextromethorphan HBr/ Guaifenesin / Phenylephrine HCL

COUGH SUPPRESSANT
EXPECTORANT/
NASAL DECONGESTANT

NON-DROWSY

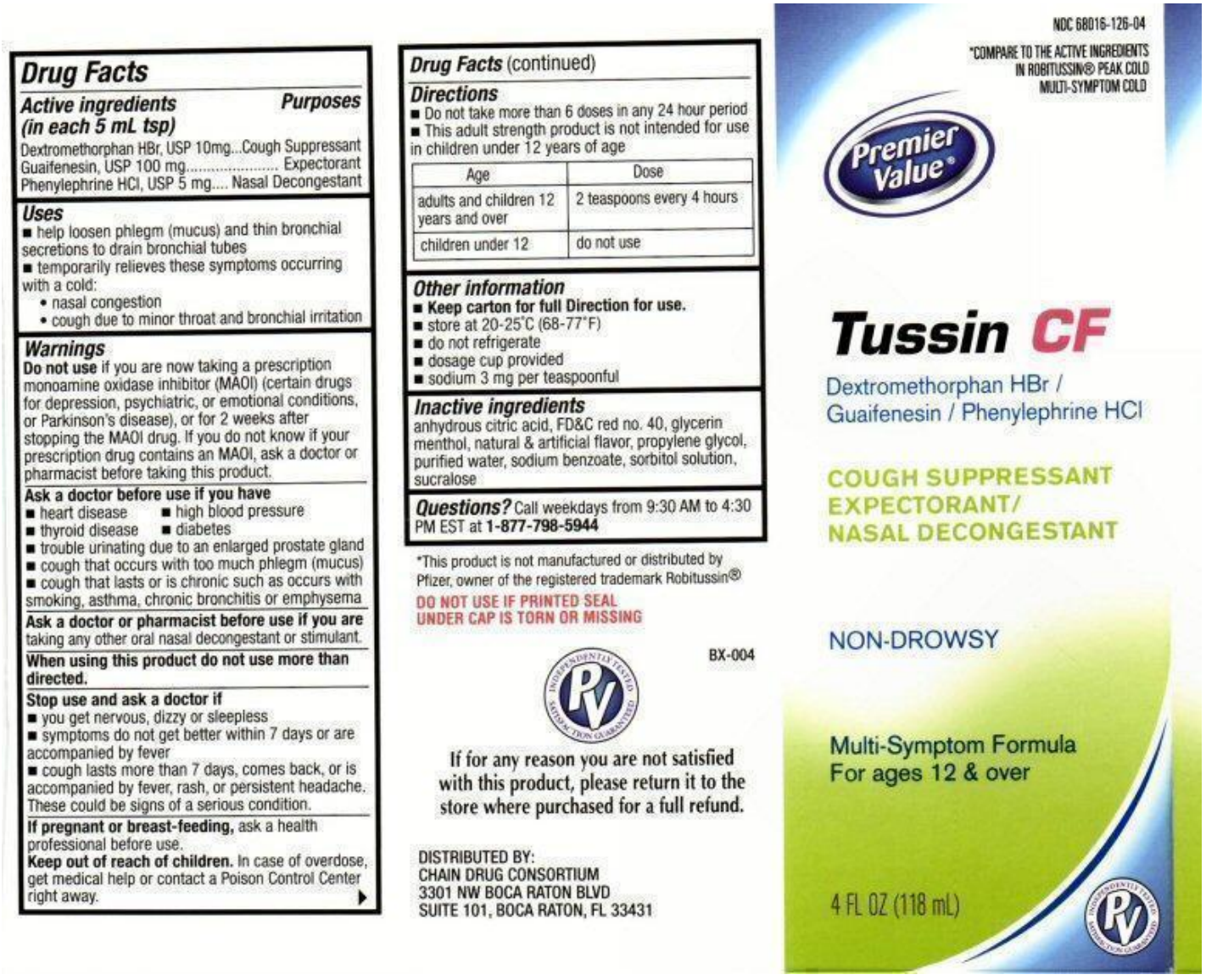
Multi-Symptom Formula

For ages 12 and over
8 FL OZ (237 mL)

INDEPENDENTLY TESTED SATISFACTION GUARANTEED PV
If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.

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

DISTRIBUTED BY: CHAIN DRUG CONSORTIUM 3301 NW BOCA RATON BLVD SUITE 101, BOCA RATON, FL 33431
BX-018
DO NOT USE IF PRINTED SEAL UNDER CAP IS TORN OR MISSING



Drug Facts

Active ingredients (in each 5 mL tsp) — **Purposes**

Dextromethorphan HBr, USP 10mg...Cough Suppressant
Guaifenesin, USP 100 mg..... Expectorant
Phenylephrine HCl, USP 5 mg.... Nasal Decongestant

Uses
- help 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

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Drug Facts (continued)

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BX-004

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DISTRIBUTED BY:
CHAIN DRUG CONSORTIUM
3301 NW BOCA RATON BLVD
SUITE 101, BOCA RATON, FL 33431

NDC 68016-126-04

*COMPARE TO THE ACTIVE INGREDIENTS IN ROBITUSSIN® PEAK COLD MULTI-SYMPTOM COLD

Premier Value®

Tussin CF

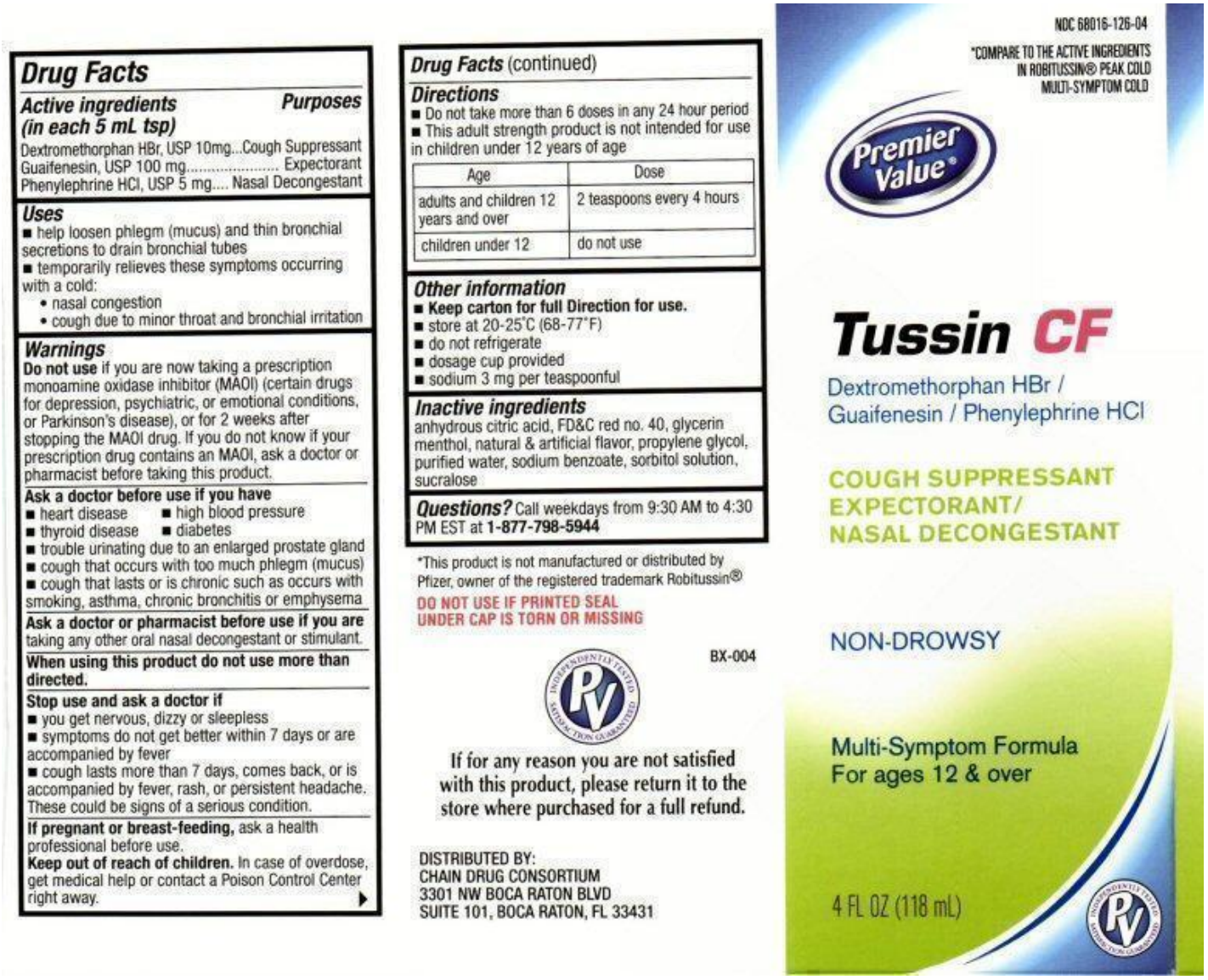
Dextromethorphan HBr / Guaifenesin / Phenylephrine HCl

COUGH SUPPRESSANT
EXPECTORANT/
NASAL DECONGESTANT

NON-DROWSY

Multi-Symptom Formula
For ages 12 & over

4 FL OZ (118 mL)



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TUSSIN CF

dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68016-126
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (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 (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	CHERRY (cherry)	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-126-04	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/15/2012	
2	NDC:68016-126-08	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/15/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	05/15/2012	

Labeler - Chain Drug Consortium, LLC (101668460)

Registrant - AptaPharma Inc. (790523323)

Establishment

Name	Address	ID/FEI	Business Operations
AptaPharma Inc.		790523323	manufacture(68016-126)

Revised: 12/2023 Chain Drug Consortium, LLC