TUSSIN DM COUGH SUPPRESSANT/EXPECTORANT- dextromethorphan hydrobromide, guaifenesin 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

Purpose

Cough Suppressant
Expectorant

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

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes

Warnings

Do not use

- in a child under 12 years of age
- 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

- 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.

Directions

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

Age	Dose
adults & children 12 years & over	2 teaspoonfuls every 4 hours
children under 12 years	do not use

Other information

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

Inactive ingredients

anhydrous citric acid, dextrose, FD and C red no.40, flavor, glycerin, high fructose corn syrup, menthol, purified water, saccharin sodium, sodium benzoate

Questions?

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

1-877-798-5944

Product Label

NDC 68016-855-56

***COMPARE TO THE ACTIVE INGREDIENTS IN ROBITUSSIN® PEAK COLD COUGH and CHEST CONGESTION DM**

Premier Value®
Tussin DM

Dextromethorphan HBr
Guaifenesin

COUGH SUPPRESSANT /
EXPECTORANT

NON-DROWSY

Helps to Loosen
Chest Congestion

Cough Formula
for ages 12 and over

8 FL OZ (237 mL)

INDEPENDENTLY TESTED SATISFACTION GUARANTEED
DO NOT USE IF IMPRINTED SHRINK BAND IS MISSING OR BROKEN

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

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

LR-018

LOT: EXP:

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

Drug Facts

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

Dextromethorphan HBr, USP 10 mg ..... Cough Suppressant
Guaifenesin, USP 100 mg ..... Expectorant

Uses

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- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes

Warnings

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■ 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
■ 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 right away.

Directions

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

Age	Dose
adults & children 12 years & over	2 teaspoonfuls every 4 hours
children under 12 years	do not use

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

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NDC 68016-855-56

*COMPARE TO THE ACTIVE INGREDIENTS IN ROBITUSSIN® PEAK COLD COUGH & CHEST CONGESTION DM

Premier Value®

Tussin DM

Dextromethorphan HBr/
Guaifenesin

COUGH SUPPRESSANT/ EXPECTORANT

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

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

BX-017

NDC 68016-855-56

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Premier Value®

Tussin DM

Dextromethorphan HBr/
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COUGH SUPPRESSANT/ EXPECTORANT

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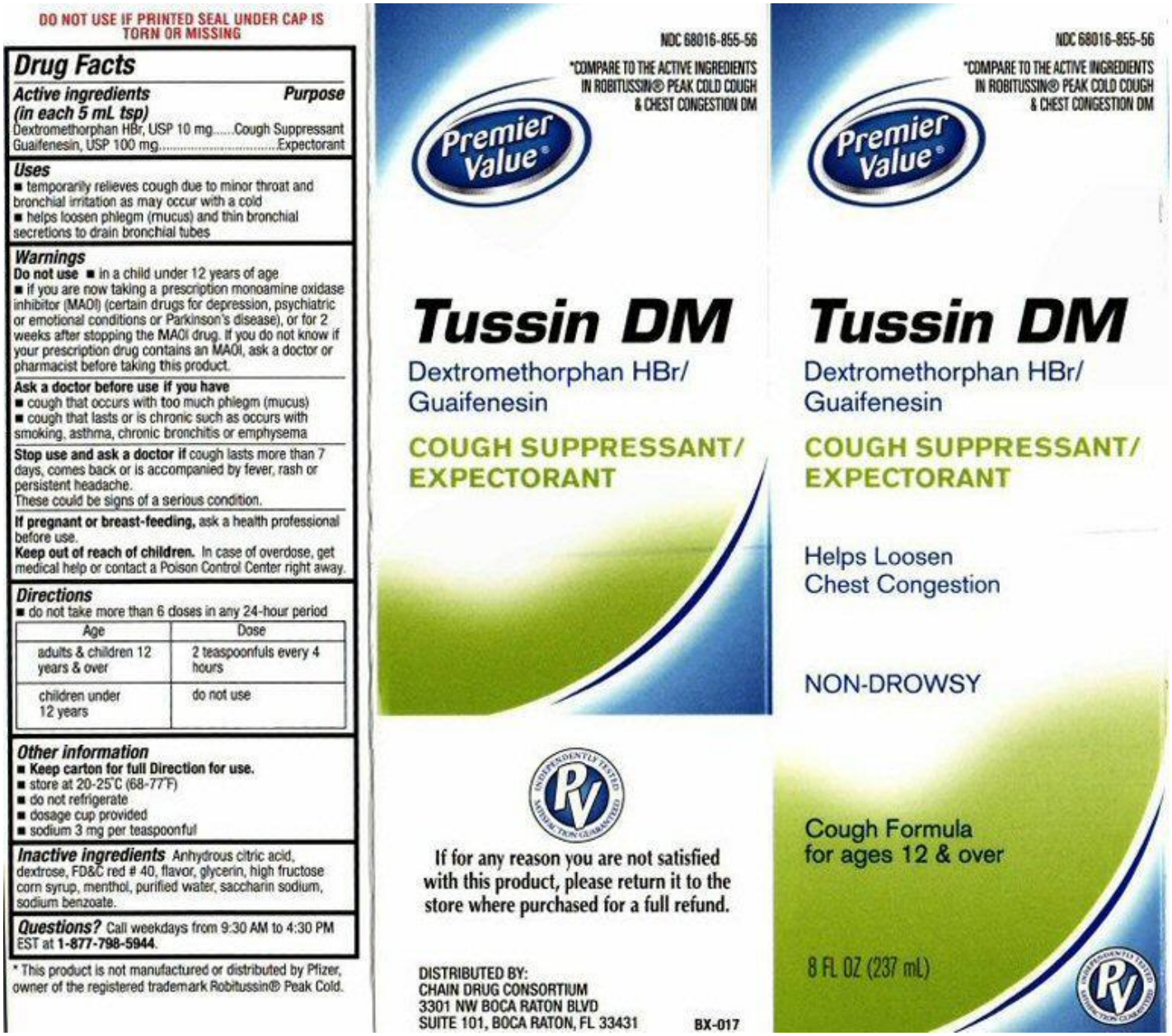

TUSSIN DM COUGH SUPPRESSANT/EXPECTORANT

dextromethorphan hydrobromide, guaifenesin liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68016-855
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

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-855-56	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2012	
2	NDC:68016-855-57	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	09/01/2012	

Labeler - Chain Drug Consortium, LLC (101668460)

Registrant - AptaPharma Inc. (790523323)

Establishment

Name	Address	ID/FEI	Business Operations
AptaPharma Inc.		790523323	manufacture(68016-855)

Revised: 12/2023 Chain Drug Consortium, LLC