**SEVERE CONGESTION AND COUGH MAX MAXIUM STRENGTH-**
**dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride liquid**
**Chain Drug Consortium, LLC**

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**Active ingredients**

**Drug Facts**

**Active ingredients (in each 20 mL)**

Dextromethorphan HBr 20 mg
Guaifenesin 400 mg
Phenylephrine HCL 10 mg

**Purpose**

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**

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

- temporarily relieves these symptoms occurring with a cold:
- cough due to minor throat and bronchial irritation
- nasal congestion

**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 two weeks after stopping the MAOI drug. If you do not know if your child’s prescription drug contains an MAOI, ask a doctor or a pharmacist before taking this product.

- for children under 12 years of age

**Ask a doctor before use if you have**
- heart disease
- thyroid disease
- high blood pressure
- diabetes
- trouble urinating due to an enlarged prostate gland
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema.
- cough that occurs with too much phlegm (mucus)

**When using this product**
- **do not use more than directed**

**Stop use and ask a doctor if**
- nervousness, dizziness or sleeplessness occur
- symptoms do not get better within 7 days or are accompanied by a fever
- cough 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.

**Directions**
- take only as recommended
- use dose cup
- mL = milliliter
- do not take more than 6 doses in any 24-hour period

**Other information**

each 20 mL teaspoon contains:

- sodium 10 mg
- dosage cup provided
- store between 15-30 ° C (59-86° F)
- do not refrigerate

**Inactive ingredients**

citric acid anhydrous, edetate disodium, FD and C Blue # 1 FD and C Red # 40, flavors, glycerin, propylene glycol, propyl gallate, purified water, sodium benzoate, sorbitol,

sucralose, Xanthan gum

**Questions?**

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

**1-877-798-5944**

**Product Label**

**NDC 68016-221-00**
***COMPARE TO THE ACTIVE INGREDIENTS IN MUCINEX® FAST-MAX™ SEVERE CONGESTION AND COUGH LF-004**

**Premier Value®**

**Maximum Strength**
**Severe Congestion and Cough MAX**

Dextromethorphan HBr / Guaifenesin / Phenylephrine HCL
COUGH SUPPRESSANT / EXPECTORANT/ NASAL DECONGESTANT

- Controls Cough
- Thins and Loosens Mucus
- Relieves Nasal and Chest Congestion

Adults

For Ages 12 and Over

6 FL OZ (177mL)

INDEPENDENTLY TESTED SATISFACTION GUARANTEED PV
**DO NOT USE IF PRINTED SEAL UNDER CAP IS TORN OR MISSING**

Peel Corner to read Complete Drug Facts and Information

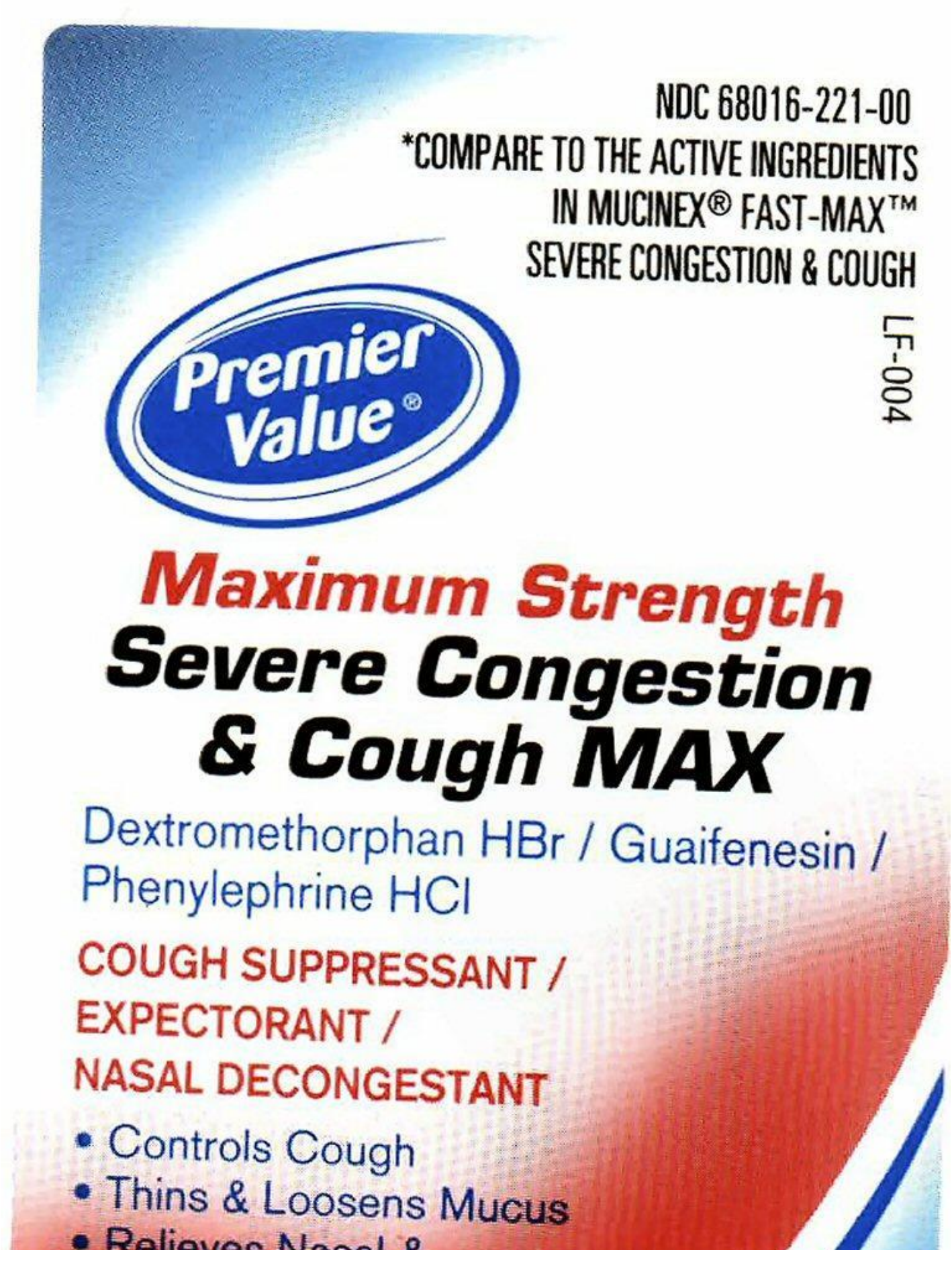
Lot: Exp:

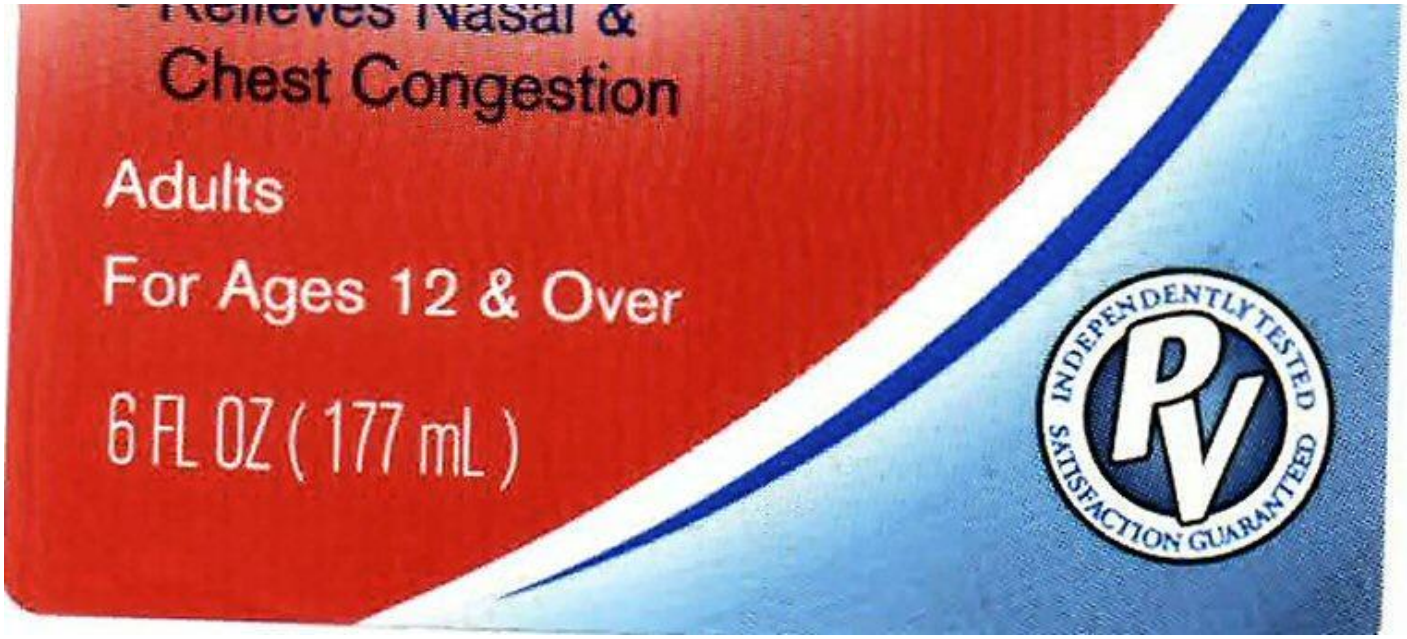
*This product is not manufactured or distributed by Reckitt Benckiser Inc. distributor of Mucinex® FAST-MAX™ DM MAX

If for Any reason you are not satisfied with this product, lease 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

LF-004


NDC 68016-221-00
*COMPARE TO THE ACTIVE INGREDIENTS IN MUCINEX® FAST-MAX™ SEVERE CONGESTION & COUGH
LF-004
Premier Value®
Maximum Strength
Severe Congestion & Cough MAX
Dextromethorphan HBr / Guaifenesin / Phenylephrine HCl
COUGH SUPPRESSANT / EXPECTORANT / NASAL DECONGESTANT
• Controls Cough
• Thins & Loosens Mucus




## Drug Facts (continued)

**If pregnant or breast feeding,** ask a health professional before use.

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## Directions

- take only as recommended
- use dosage cup
- mL = milliliter
- do not take more than 6 doses in any 24-hour period

| Age | Dose |
|---|---|
| Adults & children 12 years and older | 20 mL every 4 hours |
| Children under 12 years of age | Do not use |

## Other information

- each 20 mL contains: sodium 10 mg
- dosage cup provided
- store between 15-30°C (59-86°F)
- do not refrigerate ▸

***Peel Corner to Read Complete Drug Facts and Information***

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

LB-004

8 40986 02891 5

***Lot:*** ***Exp:***

## Drug Facts (continued)

### Inactive ingredients

anhydrous citric acid, edetate disodium, FD&C Blue #1, FD&C Red #40, flavors, glycerin, propylene glycol, propyl gallate, purified water, sodium benzoate, sorbitol, sucralose, Xanthan gum

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

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Guaifenesin 400 mg.....Expectorant

Phenylephrine HCl 10 mg..... .....Nasal Decongestant

### Uses

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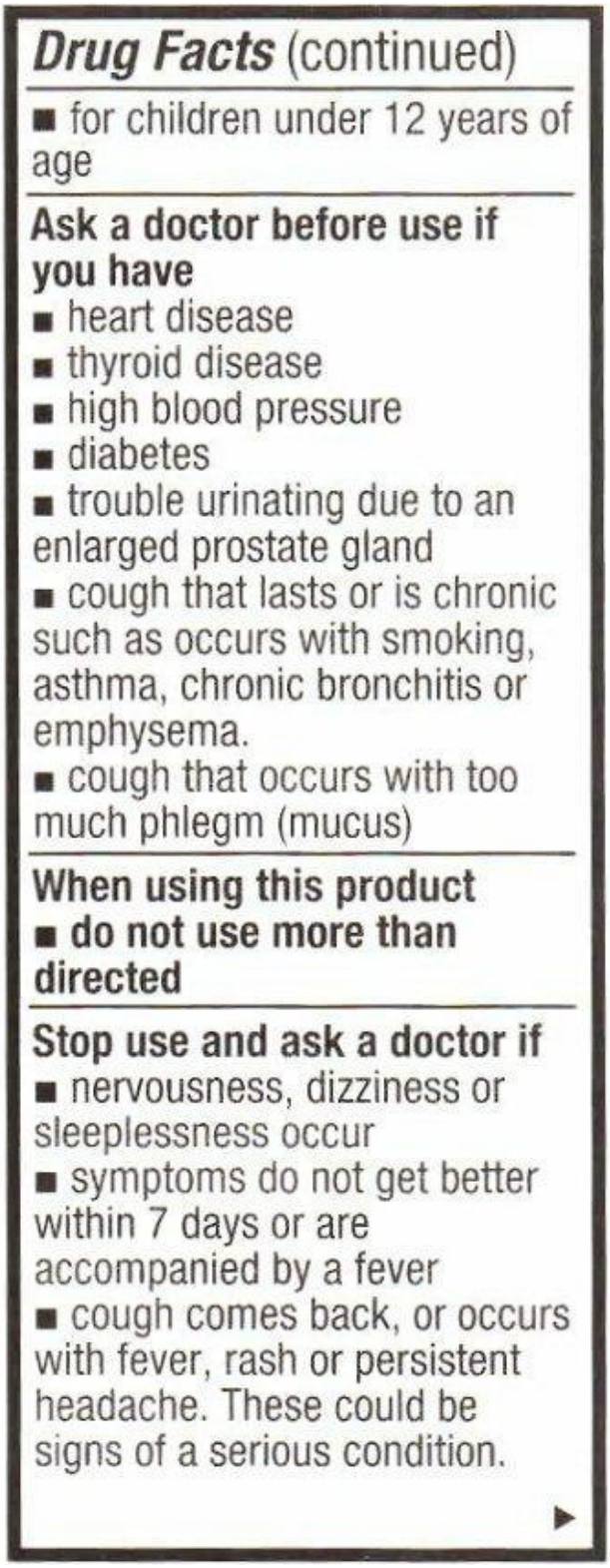

## Drug Facts (continued)

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**SEVERE CONGESTION AND COUGH MAX MAXIUM STRENGTH**

dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:68016-221 |
| **Route of Administration** | ORAL | | |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **DEXTROMETHORPHAN HYDROBROMIDE** (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN HYDROBROMIDE | 20 mg in 20 mL |
| **GUAIFENESIN** (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) | GUAIFENESIN | 400 mg in 20 mL |
| **PHENYLEPHRINE HYDROCHLORIDE** (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV) | PHENYLEPHRINE HYDROCHLORIDE | 10 mg in 20 mL |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **ANHYDROUS CITRIC ACID** (UNII: XF417D3PSL) | |
| **EDETATE DISODIUM** (UNII: 7FLD91C86K) | |
| **FD&C BLUE NO. 1** (UNII: H3R47K3TBD) | |
| **FD&C RED NO. 40** (UNII: WZB9127XOA) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **PROPYL GALLATE** (UNII: 8D4SNN7V92) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **SODIUM BENZOATE** (UNII: OJ245FE5EU) | |
| **SORBITOL** (UNII: 506T60A25R) | |
| **SUCRALOSE** (UNII: 96K6UQ3ZD4) | |
| **XANTHAN GUM** (UNII: TTV12P4NEE) | |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:68016-221-00 | 177 mL in 1 BOTTLE; Type 0: Not a Combination Product | 01/01/2013 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M012 | 01/01/2013 | |

## Labeler - Chain Drug Consortium, LLC (101668460)

## Registrant - AptaPharma Inc. (790523323)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| AptaPharma Inc. | | 790523323 | manufacture(68016-221) |