**SEVERE CONGESTION AND COUGH RELIEF MAXIMUM STRENGTH-**
**dextromethorphan hbr, guaifenesin, phenylephrine hcl solution**
**L.N.K. International, Inc.**

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**Quality Plus 44-004**

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

Dextromethorphan HBr 20 mg
Guaifenesin 400 mg
Phenylephrine HCl 10 mg

## *Purpose*

Cough suppressant
Expectorant
Nasal decongestant

## *Uses*

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- temporarily relieves:
- cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
- the intensity of coughing
- the impulse to cough to help you get to sleep
- nasal congestion due to a cold

## *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
- diabetes
- high blood pressure
- thyroid disease
- difficulty in urination due to enlargement of the prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis or emphysema
- cough that occurs with too much phlegm (mucus)

**When using this product**

**do not exceed recommended dosage.**

**Stop use and ask a doctor if**

- nervousness, dizziness, or sleeplessness occur
- symptoms do not get better within 7 days or occur with fever
- cough persists more than 7 days, tends to recur, or is accompanied by a 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 directed**
- do not take more than 6 doses in any 24-hour period
- mL = milliliter
- only use the dose cup provided
- adults and children 12 years and over: 20 mL in dosing cup provided every 4 hours
- children under 12 years: do not use

***Other information***

- **each 20 mL contains:** sodium 9 mg
- use by expiration date on package
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)

***Inactive ingredients***

anhydrous citric acid, FD&C blue #1, FD&C red #40, flavors, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate dihydrate, sodium metabisulfite, sorbitol, sucralose, xanthan gum

***Questions or comments?***

**1-888-423-0139**

***Principal Display Panel***

**QUALITY**
**+PLUS**

NDC 50844-004-45

*Compare to active ingredients
in Mucinex® FAST-MAX®
Severe Congestion & Cough

MAXIMUM
STRENGTH

**SEVERE
CONGESTION &
COUGH RELIEF**

Dextromethorphan HBr
Guaifenesin
Phenylephrine HCl

Cough Suppressant
Expectorant
Nasal Decongestant

Ages 12 Years and Over

Mixed
Berry
Flavored

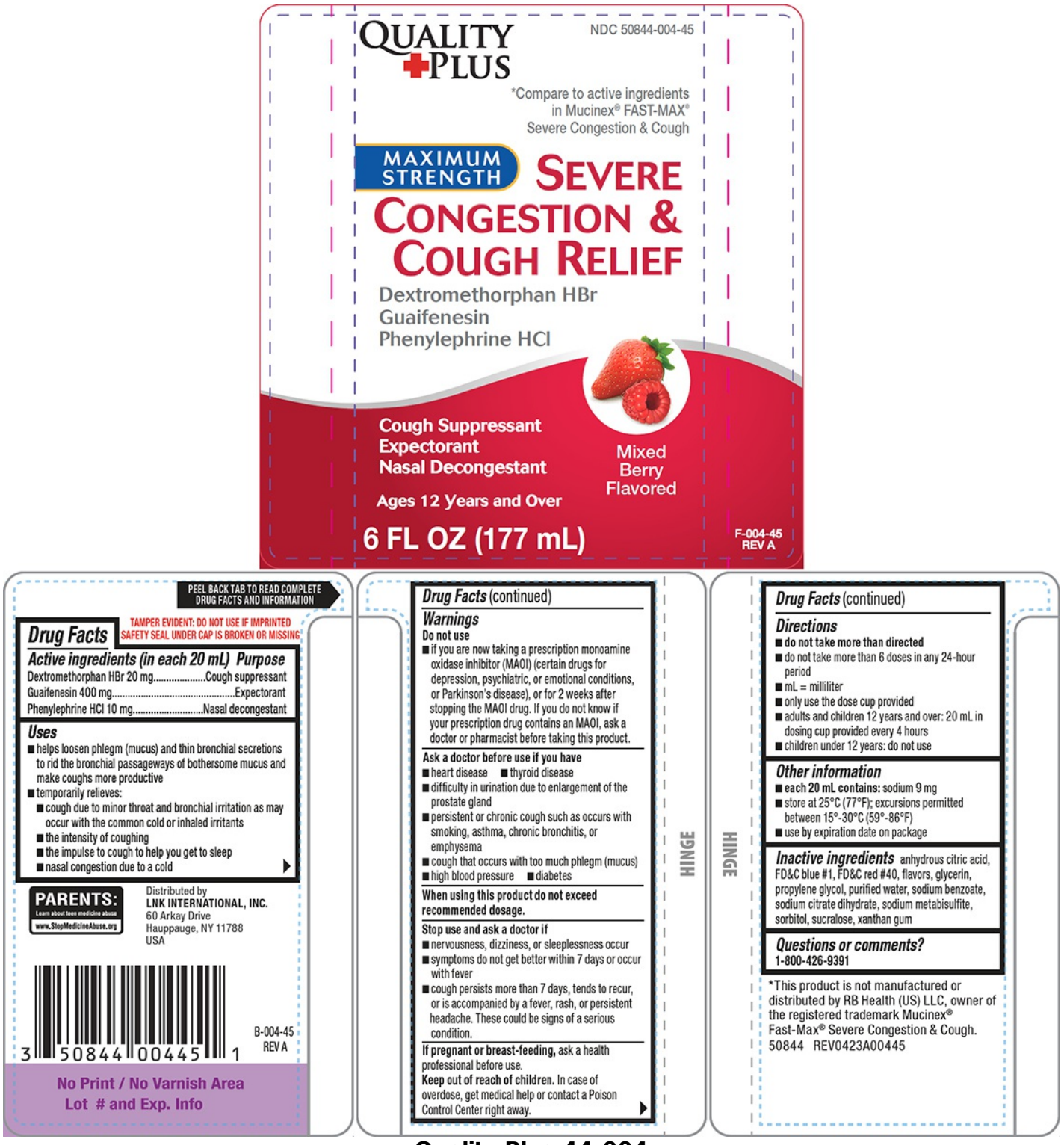
6 FL OZ (177 mL)

**TAMPER EVIDENT: DO NOT USE IF IMPRINTED
SAFETY SEAL UNDER CAP IS BROKEN OR MISSING**

**PARENTS:**
Learn about teen medicine abuse
**www.StopMedicineAbuse.org**

*This product is not manufactured or
distributed by RB Health (US) LLC, owner of
the registered trademark Mucinex®
Fast-Max® Severe Congestion & Cough.
50844 REV0423A00445

Distributed by
**LNK INTERNATIONAL, INC.**
60 Arkay Drive
Hauppauge, NY 11788
USA




**Quality Plus 44-004**

# SEVERE CONGESTION AND COUGH RELIEF MAXIMUM STRENGTH

dextromethorphan hbr, guaifenesin, phenylephrine hcl solution

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:50844-004 |
| **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) | |
| **FD&C BLUE NO. 1** (UNII: H3R47K3TBD) | |
| **FD&C RED NO. 40** (UNII: WZB9127XOA) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **SODIUM BENZOATE** (UNII: OJ245FE5EU) | |
| **TRISODIUM CITRATE DIHYDRATE** (UNII: B22547B95K) | |
| **SODIUM METABISULFITE** (UNII: 4VON5FNS3C) | |
| **SORBITOL** (UNII: 506T60A25R) | |
| **SUCRALOSE** (UNII: 96K6UQ3ZD4) | |
| **XANTHAN GUM** (UNII: TTV12P4NEE) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | blue | **Score** | |
| **Shape** | | **Size** | |
| **Flavor** | BERRY (MIXED) | **Imprint Code** | |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:50844-004-45 | 177 mL in 1 BOTTLE; Type 0: Not a Combination Product | 03/01/2017 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M012 | 03/01/2017 | |

## Labeler - L.N.K. International, Inc. (038154464)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| LNK International, Inc. | | 967626305 | manufacture(50844-004) , pack(50844-004) |

Revised: 8/2023

L.N.K. International, Inc.