**TENALIF MULTI SYMPTOM RELIEF- acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl syrup**
**OPMX LLC**

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**Tenalif Multi Symptom Relief**

***Drug Facts***

***Active ingredients***

**Purpose**

| ***Active ingredients (in each 20 mL)*** | ***Purpose*** |
|---|---|
| Acetaminophen 500 mg | Pain reliever/fever reducer |
| Dextromethorphan HBr 26.66 mg | Cough suppressant |
| Guaifenesin 400 mg | Expectorant |
| Phenylephrine HCl 10 mg | Nasal descongestant |

***Uses***

temporarily relieves these common cold/flu symptoms:

- sinus congestion & pressure
- minor aches & pains
- nasal congestion
- cough due to minor throat & bronchial irritation
- sore throat
- headache
- temporarily reduces fever
- temporarily promotes nasal and/or sinus drainage
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive.

**Warnings**

**Liver Warning:**

This product contains acetaminophen. Severe liver damage may occur if you take:
• more than 6 doses in 24 hrs, which is the maximum daily amount for this product
• with other drugs containing acetaminophen
• 3 or more alcoholic drinks every day while using this product

**Allergy Alert:**

Acetaminophen may cause severe skin reactions. Symptoms may include:
• skin reddening • blisters • rash
If a skin reaction occurs, stop use and seek medical help right away

**Sore throat warning:**

If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

**Do not use**

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- for children 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**

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

**Ask a doctor or pharmacist before use if you are**

- taking the blood thinning drug warfarin
- taking sodium restricted diet

**When using this product**

- do not exceed recommended dose (see overdose warning)

**Stop use and ask a doctor if**

- nervousness, dizziness or sleeplessness occur
- pain, nasal congestion or cough gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- cough comes back or occurs with rash or headache that lasts.

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.**

**Overdose warning**

Taking more than recommended dose (overdose) may cause liver damaage. In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical for adults & for children even if you do not notice any signs or symptoms.

### *Directions*

- do not take more than directed (see Overdose warning)
- do not take more than 6 doses in any 24 hour period
- measure only with dosing cup provided
- do not use dosing cup with other products
- dose as follows or as directed by a doctor
- mL = milliliter
- tsp = teaspoonful

| Age | Dose |
|---|---|
| adults and children 12 years of age and older | 20 mL in dosing cup provided every 4 hours |
| Children under 12 years of age | do not use |

### *Other information*

- **each 20 mL contains**
- sodium 20 mg
- Store between 20-25°C (68-77°F)
- do not refrigerate
- keep carton for complete Drug facts

### *Inactive ingredients*

Aloe vera, citric acid, disodium EDTA, FD&C red no. 40, hydroxyethyl cellulose, natural & artificial strawberry flavor, propylene glycol, purified water, sodium benzoate, sorbitol solution, sucralose

### *Questions?*

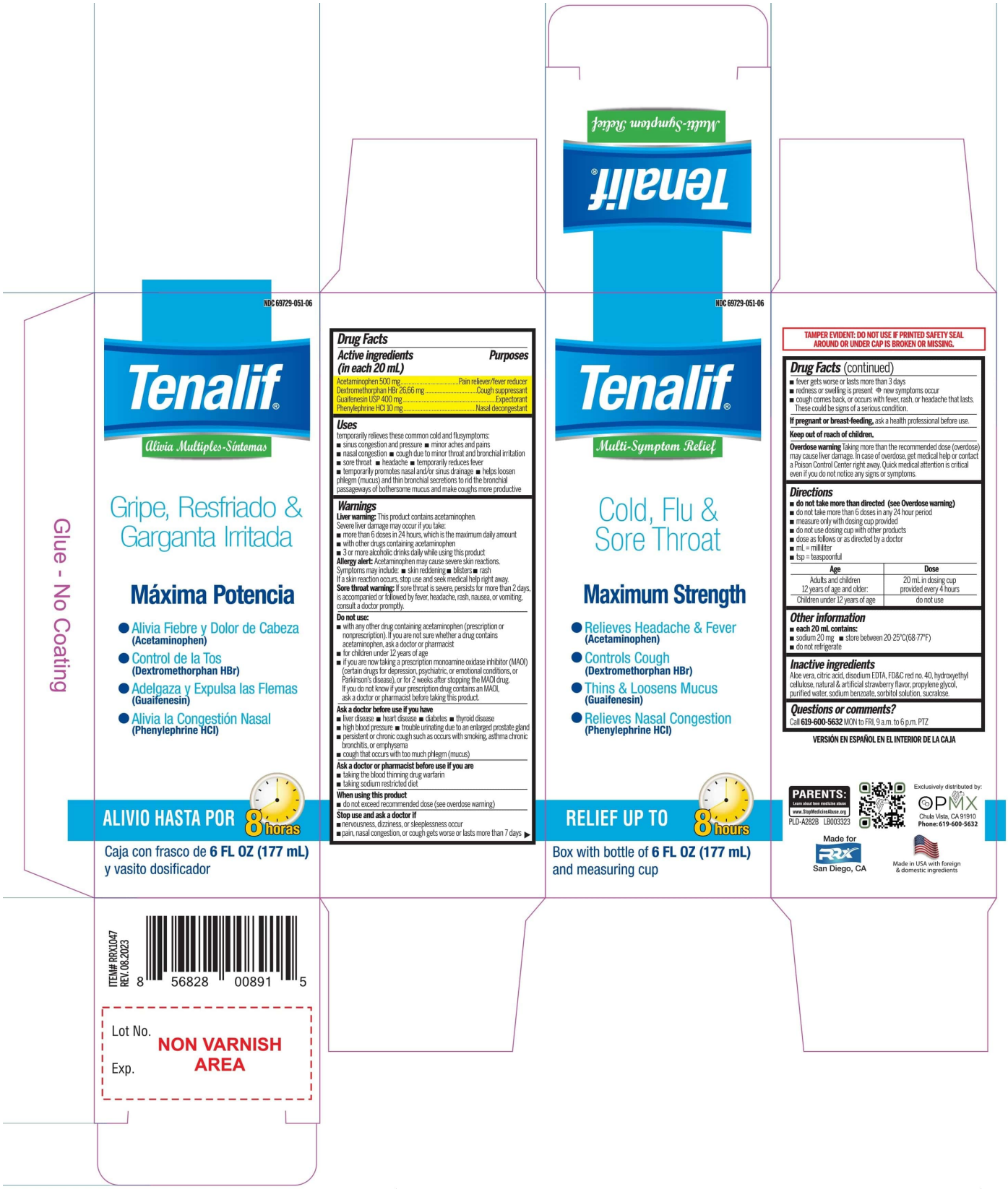
Call 616-600-5632 MON to FRI, 9 a.m. to 6 p.m. PTZ

### PRINCIPAL DISPLAY PANEL - 177 mL Bottle Label

NDC 69729-051-06

Tenalif Multi Symptom Relief

6 fl oz (177 mL)




# TENALIF MULTI SYMPTOM RELIEF

acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl syrup

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:69729-051 |

| | |
|---|---|
| **Route of Administration** | ORAL |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **ACETAMINOPHEN** (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D) | ACETAMINOPHEN | 500 mg in 20 mL |
| **PHENYLEPHRINE HYDROCHLORIDE** (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV) | PHENYLEPHRINE HYDROCHLORIDE | 10 mg in 20 mL |
| **DEXTROMETHORPHAN HYDROBROMIDE** (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN HYDROBROMIDE | 26.66 mg in 20 mL |
| **GUAIFENESIN** (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) | GUAIFENESIN | 400 mg in 20 mL |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **EDETATE DISODIUM** (UNII: 7FLD91C86K) | |
| **HYDROXYETHYL CELLULOSE, UNSPECIFIED** (UNII: T4V6TWG28D) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **ANHYDROUS CITRIC ACID** (UNII: XF417D3PSL) | |
| **ALOE** (UNII: V5VD430YW9) | |
| **SORBITOL SOLUTION** (UNII: 8KW3E207O2) | |
| **FD&C RED NO. 40** (UNII: WZB9127XOA) | |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |
| **SODIUM BENZOATE** (UNII: OJ245FE5EU) | |
| **SUCRALOSE** (UNII: 96K6UQ3ZD4) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | red | **Score** | |
| **Shape** | | **Size** | |
| **Flavor** | STRAWBERRY | **Imprint Code** | |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:69729-051-06 | 1 in 1 CARTON | 09/18/2023 | |
| **1** | | 177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M012 | 09/18/2023 | |

## Labeler - OPMX LLC (029918743)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| GADAL Laboratories, Inc | | 841305639 | manufacture(69729-051) |

Revised: 12/2024 OPMX LLC