# SANATOS TURBO MAX STRENGTH- dextromethorphan hbr, guaifenesin, phenylephrine hcl liquid Pharmadel LLC

-----

SanaTos Turbo MS (Apta)

### **Drug Facts**

#### **Active ingredients & Purposes**

| Active ingredient (in each 20 mL) | Purpose            |
|-----------------------------------|--------------------|
| Dextromethorphan HBr 20 mg        | Cough suppressant  |
| Guaifenesin 400 mg                | Expectorant        |
| Phenylephrine HCl 10 mg           | Nasal decongestant |

#### Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive
- temporarily relieves common cold/flu symptoms:
- cough due to minor throat and bronchial irritationnasal congestion due to hay fever
- other upper respiratory allergies
- sinus congestion and pressure
- stuffy nose

# Warnings

#### Do not use

- 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 two 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
- thyroid disease
- diabetes
- high blood pressure
- a cough that occurs with too much phlegm (mucus)
- a persistent or chronic cough such occurs with smoking, asthma or emphysema
- trouble urinating due to an enlarged prostate gland

### When using this product

do not use more than directed

#### Stop use and ask a doctor if

- nervousness, dizziness or sleeplessness occur
- a persistent cough or symptoms do not get better within 7 days
- cough comes back, or occurs with 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 exceed recommended dosage
- 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

# Inactive ingredients

anhydrous citric acid, dextrose, D&C red # 33, FD&C red #40, flavors, glycerin, maltitol, propylene glycol, saccharin sodium, sodium benzoate, sucralose, xanthan gum, water

#### Questions or comments?

**+1-866-359-3478** (M-F) 9 AM to 5 PM Eastern or www.pharmadel.com

\* This product is not manufactured or distributed by Reckitt Benckiser Inc., distributor of Mucinex® FAST-MAXTM Severe Congestion & Cough

#### PACKAGE PRINCIPAL DISPLAY PANEL



# **SANATOS TURBO MAX STRENGTH**

dextromethorphan hbr, guaifenesin, phenylephrine hcl liquid

| Product Information     |                |                    |               |
|-------------------------|----------------|--------------------|---------------|
| Product Type            | HUMAN OTC DRUG | Item Code (Source) | NDC:55758-322 |
| Route of Administration | ORAL           |                    |               |

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

| Inactive Ingredients               |          |  |
|------------------------------------|----------|--|
| Ingredient Name                    | Strength |  |
| PROPYL GALLATE (UNII: 8D4SNN7V92)  |          |  |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) |          |  |

| ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)  SORBITOL (UNII: 506T60A25R)  GLYCERIN (UNII: PDC6A3C0OX)  FD&C RED NO. 40 (UNII: WZB9127XOA)  EDETATE DISODIUM (UNII: 7FLD91C86K)  PROPYLENE GLYCOL (UNII: 6DC9Q167V3)  SUCRALOSE (UNII: 96K6UQ3Z D4)  SODIUM BENZOATE (UNII: OJ245FE5EU)  XANTHAN GUM (UNII: TTV12P4NEE) | WATER (UNII: 059QF0KO0R)                 |  |
|---|--|--|
| GLYCERIN (UNII: PDC6A3C0OX)  FD&C RED NO. 40 (UNII: WZB9127XOA)  EDETATE DISODIUM (UNII: 7FLD91C86K)  PROPYLENE GLYCOL (UNII: 6DC9Q167V3)  SUCRALOSE (UNII: 96K6UQ3ZD4)  SODIUM BENZOATE (UNII: OJ245FE5EU)   | ANHYDROUS CITRIC ACID (UNII: XF417D3PSL) |  |
| FD&C RED NO. 40 (UNII: WZB9127XOA)  EDETATE DISODIUM (UNII: 7FLD91C86K)  PROPYLENE GLYCOL (UNII: 6DC9Q167V3)  SUCRALOSE (UNII: 96K6UQ3ZD4)  SODIUM BENZOATE (UNII: OJ245FE5EU)  | SORBITOL (UNII: 506T60A25R)              |  |
| EDETATE DISODIUM (UNII: 7FLD91C86K)  PROPYLENE GLYCOL (UNII: 6DC9Q167V3)  SUCRALOSE (UNII: 96K6UQ3Z D4)  SODIUM BENZOATE (UNII: OJ245FE5EU)   | GLYCERIN (UNII: PDC6A3C0OX)              |  |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3)  SUCRALOSE (UNII: 96K6UQ3Z D4)  SODIUM BENZOATE (UNII: OJ245FE5EU)  | FD&C RED NO. 40 (UNII: WZB9127XOA)       |  |
| SUCRALOSE (UNII: 96K6UQ3ZD4) SODIUM BENZOATE (UNII: OJ245FE5EU)   | EDETATE DISODIUM (UNII: 7FLD91C86K)      |  |
| SODIUM BENZOATE (UNII: OJ245FE5EU)  | PROPYLENE GLYCOL (UNII: 6DC9Q167V3)      |  |
| ·   | SUCRALOSE (UNII: 96K6UQ3ZD4)             |  |
| XANTHAN GUM (UNII: TTV12P4NEE)  | SODIUM BENZOATE (UNII: OJ245FE5EU)       |  |
|   | XANTHAN GUM (UNII: TTV12P4NEE)           |  |

| l | Packaging              |  |                         |                       |
|---|------------------------|--|-------------------------|-----------------------|
|   | # Item Code            | Package Description  | Marketing Start<br>Date | Marketing End<br>Date |
|   | 1 NDC:55758-<br>322-06 | 177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 09/03/2020              |                       |

| Marketing Information |   |                         |                       |
|-----------------------|---|-------------------------|-----------------------|
| Marketing<br>Category | Application Number or Monograph<br>Citation | Marketing Start<br>Date | Marketing End<br>Date |
| OTC Monograph Drug    | M012  | 09/03/2020              |                       |
|                       |   |                         |                       |

# Labeler - Pharmadel LLC (030129680)

Revised: 11/2024 Pharmadel LLC