SANATOS MULTI SYMPTOM DAYTIME- acetaminophen, dextromethorphan hbr, phenylephrine hci liquid Pharmadel LLC

Sanatos Daytime MS 6 (APTA)

Drug Facts

Active Ingredients & Purposes

Active ingredients (in each 15 mL, 1 tablespoon) Acetaminophen 325 mg

Dextromethorphan HBr 10 mg Phenylephrine HCI 5 mg

Purposes

Pain reliever/fever reducer Cough suppressant Nasal decongestant

Uses

temporarily relieves common cold/flu symptoms:

- minor aches & pains
- headache
- sore throat
- fever
- nasal congestion
- cough due to minor throat and bronchial irritation

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if

- adults take more than 4,000 mg if acetaminophen in 24 hours
- child take more than 5 doses in 24 hours
- taken with other drugs containing acetaminophen
- adult has 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

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

Ask a doctor before use if you have

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

When using this product

- do not use more than directed (see overdose warning)
- avoid alcoholic drinks

Stop use and ask a doctor if

- you get nervos, dizzy or sleepless
- new symptoms occur
- fever gets worse or lasts more than 3 days
- pain or cough gets worse or lasts more than 5 days (children) or 7 days (adults)
- redness or swelling is present
- 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 this and all drugs out of reach of children.

Overdose warning:

In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

- take only as recommended (see overdose warning)
- use a dosage cup or tablespoon (TBSP)
- do not exceed 4 doses per 24 hrs

age	dose
adults & children 12 years & over	30 mL (2 TBSP) every 4 hours
children 6 years to under 12 years	15 mL (1 b TBSP) every 4 hours
children 4 years to under 6 years	ask a doctor
children under 4 years	do not use

 when using daytime and nighttime products, carefully read each label to ensure correct dosing.

Other information

- sodium content per tablespoon: 10 mg
- store at room temperature

TAMPER EVIDENT: Do not use if imprinted band is missing or broken.

Inactive ingredients

citric acid, FD&C yellow #6, flavor, glycerin, propylene glycol, purfied water, saccarhin sodium, sodium benzoate, sucrose

Questions?

call **1-866-359-3478**

Distributed by:

Pharmadel LLC

Seaford, DE 19973

www.sanatos.com

Principal Display Panel



SANATOS MULTI SYMPTOM DAYTIME

acetaminophen, dextromethorphan hbr, phenylephrine hci liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:55758-055

Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength **DEXTROMETHORPHAN HYDROBROMIDE** (UNII: 9D2RTI9KYH) **DEXTROMETHORPHAN** 10 mg (DEXTROMETHORPHAN - UNII:7355X3ROTS) **HYDROBROMIDE** in 15 mL PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE -**PHENYLEPHRINE** 5 mg UNII:1WS297W6MV) **HYDROCHLORIDE** in 15 mL 325 mg ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII: 36209ITL9D) | ACETAMINOPHEN in 15 mL

Inactive Ingredients		
Ingredient Name	Strength	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)		
WATER (UNII: 059QF0KO0R)		
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)		
GLYCERIN (UNII: PDC6A3C0OX)		
SUCROSE (UNII: C151H8M554)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SACCHARIN SODIUM (UNII: SB8ZUX40TY)		

Product Characteristics

Color	orange	Score
Shape		Size
Flavor	ORANGE	Imprint Code
Contains		

l	P	Packaging Packaging					
	#	Item Code Package Description		Marketing Start Date	Marketing End Date		
	1	NDC:55758- 055-06	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/01/2014	12/01/2014		
	2	NDC:55758- 055-25	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/01/2014			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M013	12/01/2014		

Labeler - Pharmadel LLC (030129680)

Revised: 11/2023 Pharmadel LLC