

MUCOSAN MAX WITH ACETAMINOPHEN- acetaminophen, dextromethorphan hydrobromide, guaifenesin and phenylephrine hydrochloride liquid
Menper Distributors, Inc.

Mucosan Max with Acetaminophen

Active Ingredients (in each 20 mL) Purpose

Acetaminophen, 650 mg.....Pain reliever/Fever reducer

Dextromethorphan HBr, 20 mg.....Cough suppressant

Guaifenesin, 400 mg.....Expectorant

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

Pain reliever/Fever reducer

Cough suppressant

Expectorant

Nasal Decongestant

Uses

- temporarily relieves these common cold and flu symptoms:
- nasal congestion
- cough
- minor aches and pains
- sore throat
- headache
- temporarily reduces fever
- helps loosen phlem (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 4 doses (20 mL each) in 24 hours, wich is the maximum daily amount for this product
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks daily while using this product.

Allergy alert:acetaminophen may cause severe skin reactions. Symptoms may include: skin reddening, blisters and rash. If 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 acetamonophen (prescription or nonprescription). If

you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist

- if you are 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 a MAOI, ask a doctor or Pharmacist before taking this product.

Ask a doctor before use if you have

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

Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin.

When using this product do not use more than direct.

Stop use and ask a doctor if

- you get nervous, dizzy or sleepless
- symptoms do not get better within 7 days or are accompanied by fever
- cough comes back or occurs with fever, rash, or persistant headache
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur. 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 the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical even if you do not notice any signs or symptoms.

Directions

- **do not take more than direct**
- do not take more than 4 doses in any 24 hours
- this adult strength product is not intended for use in children under 12 years of age
- measure only with dosing cup provided
- keep dosing cup with product
- mL = milliliter
- dose as follows

age	dose
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adults and children 12 years of age and older	20 mL every 4 hours
children under 12 years of age	do not use

Other information

- **each (20 mL) contains:**sodium 10mg
- store between 15-30°C (59-86°F)
- do not refrigerate.

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 or comments?

1-800-560-5223 Monday to Friday 9am-4pm, Eastern Time



PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 20 mL
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Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
XANTHAN GUM (UNII: TTV12P4NEE)	
WATER (UNII: 059QF0KOOR)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL (UNII: 506T60A25R)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53145-696-06	1 in 1 CARTON	11/01/2024	
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	11/01/2024	

Labeler - Menper Distributors, Inc. (101947166)

Registrant - Menper Distributors, Inc. (101947166)

Revised: 1/2026

Menper Distributors, Inc.