## MAXIMUM STRENGTH MUCUS RELIEF COLD, FLU AND SORE THROATacetaminophen, dextromethorphan hydrobromide, guaifenes in , phenylephrine hcl liquid SUPERVALU INC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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# Equaline® maximum strength‡ mucus relief cold, flu and sore throat 6FL OZ

### **Drug Facts**

Active ingredients (in each 20 mL)	Purposes	
Acetaminophen 650 mg	Pain reliever/fever reducer	
Dextromethorphan HBr 20 mg	Cough suppressant	
Guaifenesin 400 mg	Expectorant	
Phenylephrine HCl 10 mg	Nasal decongestant	

#### Uses

- temporarily relieves these common cold and flu symptoms:
  - nasal congestion
  - sinus congestion and pressure
  - cough due to minor throat and bronchial irritation
  - minor aches and pains
  - 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 hours, which is the maximum daily amount
- 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
- 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
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough such as occurs with 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

## When using this product

do not use more than directed

#### 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 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.

# 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 (1-800-222-1222). Quick medical attention is critical for adults as well as for children even if you don't 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
- 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 8 mg
- store at room temperature
- do not refrigerate
- dosing cup provided

## **Inactive ingredients**

anhydrous citric acid, edetate disodium, FD&C blue #1, FD&C red #40, flavors, potassium citrate, propylene glycol, propyl gallate, purified water, sodium benzoate, sorbitol, sucralose, xanthan gum

#### Questions or comments?

1-877-932-7948

#### PRINCIPAL DISPLAY PANEL

**EQUALINE**<sup>®</sup>

NDC 41163-737-06

Compare to Mucinex® Fast-Max<sup>TM</sup> Cold, Flu & Sore Throat active ingredients\*

maximum strength‡

mucus relief

cold, flu & sore throat

**acetaminophen** (pain reliever-fever reducer) dextromethorphan HBr (cough suppressant) guaifenesin (expectorant) phenylephrine HCl ( nasal decongestant)

- relieves headache & fever
- control cough

- relieves nasal & chest congestion
- thins & loosens mucus

## For ages 12+

6FL OZ (180mL)

Tamper evident: do not use if printed seal under cap is broken or missing.

‡Maximum Strength per 4 hour dose.

**DISTRIBUTED BY** 

SUPERVALU INC,

EDEN PRAIRIE,

**MN 55344 USA** 

877-932-7948

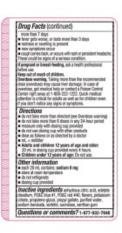
Supervaluprivatebrands.com

\*This product is not manufactured or distributed by Reckitt Benckiser, the distributor of Maximum Strength Mucinex $^{\mathbb{R}}$  Fast-Max $^{\mathsf{TM}}$  Cold, Flu & Sore Throat.









### MAXIMUM STRENGTH MUCUS RELIEF COLD, FLU AND SORE THROAT

acetaminophen, dextromethorphan hydrobromide, guaifenesin ,phenylephrine hcl liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41163-737
Route of Administration	ORAL		

# Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D)	ACETAMINOPHEN	650 mg in 20 mL
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9 D2RTI9 KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 20 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL
PHENYLEPHRINE HYDRO CHLO RIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 20 mL

Inactive Ingredients		
Ingredient Name	Strength	
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)		
EDETATE DISO DIUM (UNII: 7FLD9 1C8 6 K)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		
POTASSIUM CITRATE (UNII: EE90ONI6FF)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
PROPYL GALLATE (UNII: 8 D4SNN7V92)		
WATER (UNII: 059QF0KO0R)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SORBITOL (UNII: 506T60A25R)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		
XANTHAN GUM (UNII: TTV12P4NEE)		

Product Characteristics			
Color	BLUE	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:41163-737- 06	180 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/07/2017	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	09/07/2017	

# Labeler - SUPERVALU INC (006961411)

Revised: 9/2017 SUPERVALU INC