MUCINEX SINUS-MAX SEVERE CONGESTION RELIEF- acetaminophen, guaifenesin, and phenylephrine hydrochloride solution RB Health (US) LLC

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.

Mucinex® Sinus-Max [®] Severe Congestion Relief

Drug Facts

Active ingredients (in each 20 mL)	Purposes
Acetaminophen 650 mg	Pain reliever
Guaifenesin 400 mg	Expectorant
Phenylephrine HCl 10 mg	Nasal decongestant

Uses

- temporarily relieves:
 - nasal congestion
 - headache
 - minor aches and pains
 - sinus congestion and pressure
- temporarily promotes nasal and/or sinus drainage
- helps loosens 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.

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.
- 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 aretaking 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 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 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 for adults as well as 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
- 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 12 mg
- store between 20-25°C (68-77°F)
- dosing cup provided
- do not refrigerate

Inactive ingredients

anhydrous citric acid, edetate disodium, FD&C Blue no. 1, FD&C Red no. 40, flavors, glycerin, propyl gallate, propylene glycol, purified water, sodium benzoate, sorbitol, sucralose, trisodium citrate dihydrate ¹, xanthan gum

1 may contain this ingredient

Questions?

1-866-MUCINEX (1-866-682-4639)

You may also report side effects to this phone number.

Dist. by: Reckitt Benckiser Parsippany, NJ 07054-0224 Made in England

PRINCIPAL DISPLAY PANEL - 180 mL Bottle Label

MAXIMUM STRENGTH*

NDC 63824-261-66

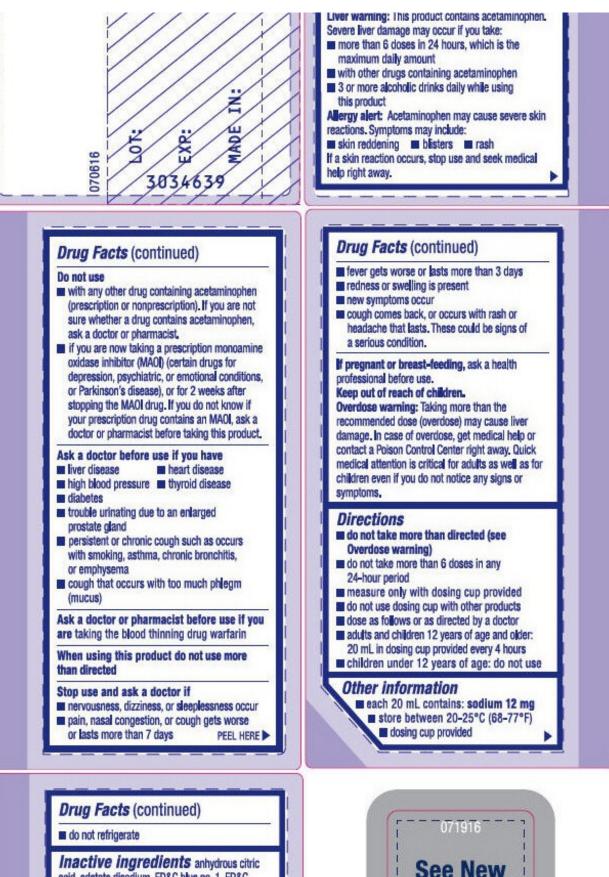
Mucinex® SINUS- MAX®

SEVERE CONGESTION RELIEF

Acetaminophen– Pain Reliever Guaifenesin – Expectorant Phenylephrine HCI – Nasal Decongestant

- ✓ Clears Sinus Congestion
- ✓ Relieves Headache
- ✓ Thins & Loosens Mucus

6 FL OZ (180mL) FOR AGES 12+ 



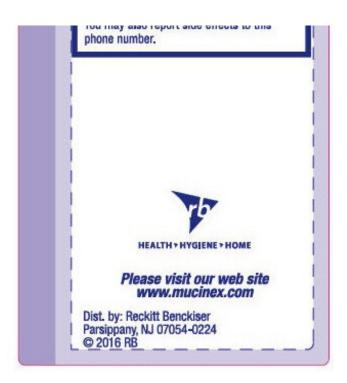
nformation

3034605

acid, edetate disodium, FD&C blue no. 1, FD&C red no. 40, flavors, glycerin, propyl gallate, propylene glycol, purified water, sodium benzoate, sorbitol, sucralose, trisodium citrate dihydrate*, xanthan gum *may contain this ingredient

Questions?

1-866-MUCINEX (1-866-682-4639) You may also report side affects to this



MUCINEX SINUS-MAX SEVERE CONGESTION RELIEF

acetaminophen, guaifenesin, and phenylephrine hydrochloride solution

Product Type	HUMAN OTC DRUG	ltem Code (Source)	NDC:63824-261
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	650 mg in 20 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 20 mL

Inactive Ingredients	
Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZ B9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

	RBITOL (UNII: 5	00100A23R)						
SU	CRALOSE (UNII:	96K6UQ3ZD4)						
TR		TE DIHYDRATE	(UNII: B22547B95	5K)				
XA	NTHAN GUM (U	NII: TTV12P4NEE)						
Pı	oduct Char	acteristics						
Со	lor		blue	Score				
Sh	аре			Size				
Fla	avor			Imprint Code				
Co	ntains							
Pa	ackaging							
			Package Des	scription		Market Start D		Marketing End Date
#	ackaging Item Code NDC:63824- 180		Type 9: Other T	- ype of Part 3 Com	bination		ate	-
#	ackaging Item Code NDC:63824- 180	mL in 1 BOTTLE;	Type 9: Other T	- ype of Part 3 Com	bination	Start D	ate	End Date
#	ackaging Item Code NDC:63824- 180	mL in 1 BOTTLE;	Type 9: Other T	- ype of Part 3 Com	bination	Start D	ate	End Date
# 1	Ackaging Item Code NDC:63824- 261-66 Pro	mL in 1 BOTTLE;	Type 9: Other T Device/Biological	- ype of Part 3 Com	bination	Start D	ate	End Date
#	Ackaging Item Code NDC:63824- 261-66 Pro) mL in 1 BOTTLE; duct (e.g., Drug/D	Type 9: Other T Device/Biological	ype of Part 3 Com Product)	bination Marketir Da	Start D 03/26/2014	ate	End Date

Labeler - RB Health (US) LLC (081049410)

Revised: 8/2023

RB Health (US) LLC