MAXIMUM STRENGTH SINUS RELIEF DAY AND NIGHT SOFTGELS - 24 CT-
acetaminophen, dextromethorphan hydrobromide, guaifenesin, doxylamine succinate, phenylephrine hydrochloride
Topco Associates LLC

MAXIMUM STRENGTH Sinus Relief Day and Night SOFTGELS**

Drug Facts

Active ingredients (in each softgel) DAY

Acetaminophen 325 mg
Dextromethorphan HBr 10 mg
Guaifenesin 200 mg
Phenylephrine HCl 5 mg

Active ingredients (in each softgel) NIGHT

Acetaminophen 325 mg
Dextromethorphan HBr 10 mg
Doxylamine succinate 6.25 mg
Phenylephrine HCl 5 mg

Purposes

Pain reliever
Cough suppressant
Expectorant
Nasal decongestant

Purposes

Pain reliever
Cough suppressant
Antihistamine
Nasal decongestant

Uses

• temporarily relieves:
  • nasal congestion • headache • cough
  • minor aches & pains • sinus congestion & pressure
  • runny nose and sneezing ***(NIGHT only)***
• 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 ***(DAY only)***

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take:
• more than 12 softgels in 24 hours, which is the maximum daily amount
• with other drugs containing acetaminophen
• 3 or more alcoholic drinks every day while using these products
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 • diabetes
• high blood pressure • thyroid disease
• trouble urinating due to an enlarged prostate gland
• glaucoma ***(NIGHT only)***
• a breathing problem such as emphysema or chronic bronchitis ***(NIGHT only)***
• 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
• taking sedatives or tranquilizers ***(NIGHT only)***

When using these products
• **do not use more than directed**
• excitability may occur, especially in children ***(NIGHT only)***
• marked drowsiness may occur ***(NIGHT only)***
• alcohol, sedatives, and tranquilizers may increase drowsiness ***(NIGHT only)***
• avoid alcoholic drinks ***(NIGHT only)***
• be careful when driving a motor vehicle or operating machinery ***(NIGHT only)***

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: In case of overdose, get medical help or contact a Poison Control

Center right away.

Directions

- **do not take more than directed (see Overdose warning)**
- do not take more than 12 softgels in any 24-hour period
- adults and children 12 years of age and older: take 2 softgels every 4 hours
- children under 12 years of age: do not use

Other information

- store at controlled room temperature between 20-25°C (68-77°F)
- protect from light, heat and moisture

Inactive ingredients (DAY only)

FD&C Yellow No. 6, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, sorbitol sorbitan solution, titanium dioxide

Inactive ingredients (NIGHT only)

D&C Yellow No.10, FD&C Blue No.1, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, sorbitol sorbitan solution, titanium dioxide

Questions or comments?

1-888-423-0139

TopCare® health

COMPARE TO MUCINEX® SINUS-MAX® DAY ACTIVE INGREDIENTS*

COMPARE TO MUCINEX® SINUS-MAX® NIGHT ACTIVE INGREDIENTS*

MAXIMUM STRENGTH**

*These products are not manufactured or distributed by Reckitt Benckiser, distributor of Mucinex® Sinus-Max® Day & Night Softgel

QUALITY GUARANTEED

This TopCare® product is laboratory tested to guarantee its highest quality. Your total satisfaction is guaranteed.

READ AND KEEP OUTER CARTON FOR COMPLETE PRODUCT WARNINGS AND INFORMATION

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PRODUCT OF INDIA

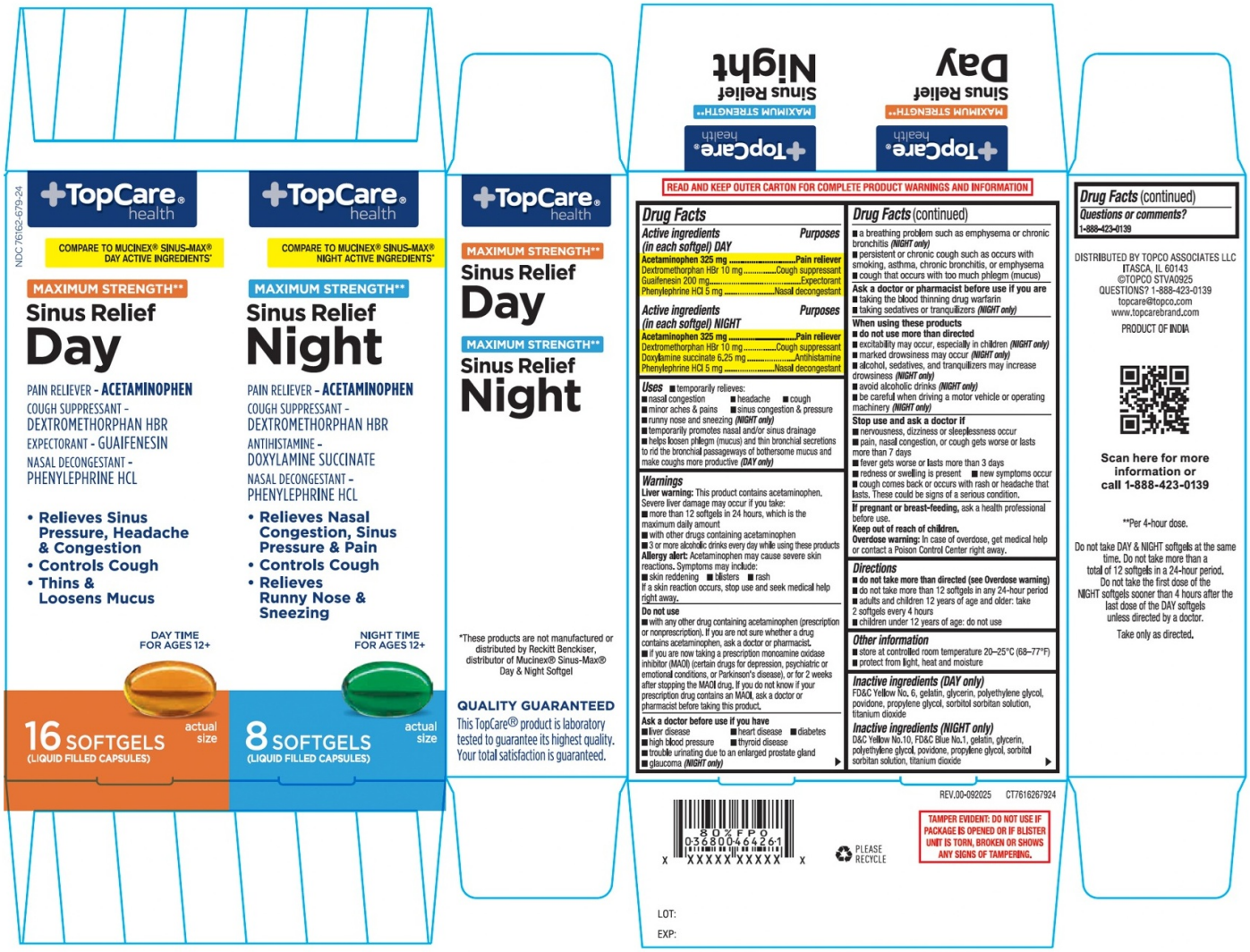
**Per 4-hour dose.

Do not take DAY & NIGHT softgels at the same time. Do not take more than a total of 12 softgels in a 24-hour period. Do not take the first dose of the NIGHT softgels sooner than 4 hours after the last dose of the DAY softgels unless directed by a doctor.

Take only as directed.

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.

Packaging



MAXIMUM STRENGTH SINUS RELIEF DAY AND NIGHT SOFTGELS - 24 CT

acetaminophen, dextromethorphan hydrobromide, guaifenesin, doxylamine succinate, phenylephrine hydrochloride kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76162-679

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76162-679-24	1 in 1 CARTON; Type 0: Not a Combination Product	10/13/2025	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	2 BLISTER PACK	16
Part 2	2 BLISTER PACK	8

Part 1 of 2

SINUS RELIEF DAY

acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride capsule, liquid filled

Product Information

Item Code (Source)	NDC:76162-680
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SORBITOL (UNII: 506T60A25R)	
SORBITAN (UNII: 6O92ICV9RU)	

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	orange	**Score**	no score
Shape	OVAL	**Size**	19mm
Flavor		**Imprint Code**	A13
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		8 in 1 BLISTER PACK; 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	10/13/2025	

Part 2 of 2

SINUS RELIEF NIGHT

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hydrochloride capsule, liquid filled

Product Information	
Item Code (Source)	NDC:76162-681
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients

Ingredient Name	Strength

D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SORBITOL (UNII: 506T60A25R)	
SORBITAN (UNII: 6O92ICV9RU)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	green	**Score**	no score
Shape	OVAL	**Size**	18mm
Flavor		**Imprint Code**	A07
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		4 in 1 BLISTER PACK; 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	10/13/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	10/13/2025	

Labeler - Topco Associates LLC (006935977)

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