TARGET MAXIMUM STRENGTH COLD FLU AND SORE THROAT OVERNIGHT COLD AND FLU- acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl and triprolidine hcl
TARGET CORPORATION

Maximum Strength Cold, Flu & Sore Throat and Overnight Cold & Flu Value Pack

Drug Facts

Active ingredients (in each 20 mL)	***Purposes***
MAXIMUM STRENGTH‡ COLD, FLU & SORE THROAT	
Acetaminophen 650 mg	**Pain reliever/fever reducer**
Dextromethorphan HBr 20 mg	Cough suppressant
Guaifenesin 400 mg	Expectorant
Phenylephrine HCl 10 mg	Nasal decongestant

Active ingredients (in each 20 mL)	***Purposes***
MAXIMUM STRENGTH OVERNIGHT COLD & FLU	
Acetaminophen 650 mg	**Pain reliever/fever reducer**
Dextromethorphan HBr 20 mg	Cough suppressant
Triprolidine HCl 2.5 mg	Antihistamine

Uses

MAXIMUM STRENGTH COLD, FLU & SORE THROAT

- temporarily relieves these common cold and flu symptoms:
- cough
- nasal congestion
- minor aches and pains
- sore throat
- headache
- stuffy nose
- sinus congestion and pressure
- temporarily reduces fever
- temporarily promotes nasal and for 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 warnings: This product contains acetaminophen. Severe liver damage may occur if you take:

- more than6 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 warnings: 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.
- 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.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children

Overdose warnings.

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 at 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 over:** 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

Inactive ingredients (Maximum strength Cold, Flu and Sore Throat)

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

Uses (OVERNIGHT COLD & FLU

- temporarily relieves these common cold and flu symptoms:
- cough
- minor aches and pains
- sore throat
- headache
- runny nose
- sneezing
- itching of the nose or throat
- itchy, watery eyes due to hay fever
- temporarily reduces fever
- controls cough to help you get to sleep

Warnings

Liver warnings: This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 4000 mg in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks daily while using this product

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 Centre right away at 1-800-222-1222.

Quick medical attention is critical for adults as well as for children, even if you do not notice any signs

Directions

- **do not take more than directed (see overdose warnings**
- do not take more than 4 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 over:** 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 10 mg
- low sodium
- store at room temperature
- do not refrigerate

Inactive ingredients (Overnight Cold & Flu)

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

Questions or comments?

(1-866-467-2748)

PRINCIPAL DISPLAY PANEL - Kit Carton

VALUE PACK

NDC 11673-691-12

Compare to active ingredients Maximum Strength‡ Mucinex® Fast Max® Cold, Flu and Sore Throat

Maximum Strength‡

Cold, Flu and Sore Throat

Acetaminophen • Pain Reliever/Fever Reducer

Dextromethorphan HBr • Cough Suppressant
Guaifenesin • Expectorant

Phenylephrine HCl • Nasal Decongestant

- **Controls Cough, Thins & Loosens Mucus**

- **Nasal & Chest Congestion**
- **Sinus pressure & Congestion**
- **Body Pain, Headache, Fever & Sore Throat**

*Per 4-hour dose

For Ages 12+

6 FL. OZ. (180 mL)

*This product is not manufactured or distributed by Reckitt Benckiser, the distributor of Maximum Strength Mucinex® Fast-Max® Cold, Flu & Sore Throat.

Compare to Mucinex® Nightshift Cold, Flu & Sore Throat Active Ingredients**

Overnight Cold & Flu

Acetaminophen • Pain Reliever/Fever Reducer
Dextromethorphan HBr • Cough Suppressant

Triprolidine HCl • Antihistamine

Night Time Relief for a Better Morning

Maximum Strength per 4-hour dose

- **Cough**
- **Fever**
- **Sore Throat**
- **Runny Nose**
- **Sneezing**

For Ages 12+

6 FL OZ (180 mL)

**This product is not manufactured or distributed by RB Health, the distributor of Mucinex® Nightshift Cold & Flu.

TAMPER EVIDENT: DO NOT USE IF PRINTED INNER SEAL UNDER CAP IS BROKEN OR MISSING.

See bottle for full labeling

Distributed by:

094 14 8715 R00

C-001227-01-055

Dist. By Target Corp.

Mpls., MN 55403

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Target Brands, Inc.

Questions?

Cal 1-866-467-2748

DO NOT TAKE MAXIMUM STRENGTH COLD, FLU & SORE THROAT & OVERNIGHT COLD & FLU LIQUIDS AT THE SAME TIME.

TARGET MAXIMUM STRENGTH COLD FLU AND SORE THROAT OVERNIGHT COLD AND FLU

acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl and triprolidine hcl kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-691

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-691-12	1 in 1 CARTON; Type 0: Not a Combination Product	03/30/2020	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	180 mL
Part 2	1 BOTTLE	180 mL

Part 1 of 2

MAXIMUM STRENGTH COLD FLU AND SORE THROAT

acetaminophen, dextromethorphan hbr, guaifenesin and phenylephrine hcl solution

Product Information	
Item Code (Source)	NDC:82442-737

Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg in 20 mL
Dextromethorphan Hydrobromide (UNII: 9D2RTI9KYH) (Dextromethorphan - UNII:7355X3ROTS)	Dextromethorphan Hydrobromide	20 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: WZB9127XOA)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
propylene glycol (UNII: 6DC9Q167V3)	
propyl gallate (UNII: 8D4SNN7V92)	
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	FRUIT	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82442-737-06	180 mL in 1 BOTTLE; Type 9: Other Type of Part 3 Combination Product (e.g., Drug/Device/Biological Product)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/30/2020	

Part 2 of 2

TARGET OVERNIGHT COLD AND FLU

acetaminophen, dextromethorphan hbr, phenylephrine hcl and triprolidine hcl solution

Product Information	
Item Code (Source)	NDC:82442-698
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Acetaminophen (UNII: 362O9ITL9D) (Acetaminophen - UNII:362O9ITL9D)	Acetaminophen	650 mg in 20 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 20 mL
TRIPROLIDINE HYDROCHLORIDE (UNII: YAN7R5L890) (TRIPROLIDINE - UNII:2L8T9S52QM)	TRIPROLIDINE HYDROCHLORIDE	2.5 mg in 20 mL

Inactive Ingredients	
Ingredient Name	**Strength**
anhydrous citric acid (UNII: XF417D3PSL)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
edetate disodium (UNII: 7FLD91C86K)	
FD&C Blue No. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
glycerin (UNII: PDC6A3C0OX)	
propyl gallate (UNII: 8D4SNN7V92)	
propylene glycol (UNII: 6DC9Q167V3)	
water (UNII: 059QF0KO0R)	
sodium benzoate (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
sorbitol (UNII: 506T60A25R)	
sucralose (UNII: 96K6UQ3ZD4)	
xanthan gum (UNII: TTV12P4NEE)	

Product Characteristics			
Color	BLUE	**Score**	
Shape		**Size**	
Flavor	FRUIT	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82442-698-06	180 mL in 1 BOTTLE; Type 9: Other Type of Part 3 Combination Product (e.g., Drug/Device/Biological Product)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/30/2020	

Labeler - TARGET CORPORATION (006961700)

Revised: 4/2024

TARGET CORPORATION