MAXIMUM STRENGTH NIGHTTIME COLD AND FLU- acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl capsule, liquid filled
TARGET CORP

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.

TARGET 11673-979 MAX STRENGTH NIGHTTIME COLD AND FLU

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

FD&C Blue No. 1, gelatin, glycerin,polyethylene glycol , povidone, propylene glycol, purified water, shellac, sorbitol sorbitan solution titanium dioxide,

Purpose
- Pain reliever/Fever reducer
- Cough suppressant
- Antihistamine
- Nasal decongestant

- temporarily relieves common cold/flu symptoms:

nasal congestion, cough, nasal congestion, minor aches and pains, sore throat, headache, runny nose and sneezing
- temporarily reduces fever
- controls cough to help you get to sleet

Directions -Do not take more than directed (see Overdose warning)
- do not exceed more than 8 softgels in any 24 hr- period -
- adults & children 12 yrs & older: take 2 Softgels every 4 hrs
- children under 12 yrs of age - Do not use.

Liver warning - This product contains acetaminophen. Severe liver damage may occur if you take:
- more than 8 Softgels in 24 hours, which 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
- Rash

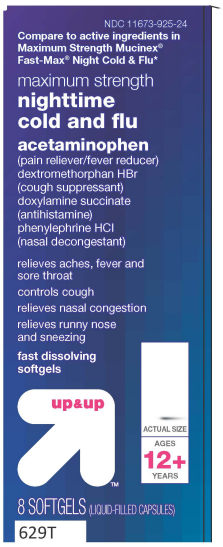
Sore troat warning

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.

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 do not notice any signs or symptoms.


NDC 11673-925-24
Compare to active ingredients in Maximum Strength Mucinex® Fast-Max® Night Cold & Flu*
maximum strength
nighttime cold and flu
acetaminophen
(pain reliever/fever reducer)
dextromethorphan HBr
(cough suppressant)
doxylamine succinate
(antihistamine)
phenylephrine HCl
(nasal decongestant)
relieves aches, fever and sore throat
controls cough
relieves nasal congestion
relieves runny nose and sneezing
fast dissolving softgels
up&up
ACTUAL SIZE
AGES
12+
YEARS
8 SOFTGELS (LIQUID-FILLED CAPSULES)
629T

MAXIMUM STRENGTH NIGHTTIME COLD AND FLU

acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl capsule, liquid filled

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-979
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
DOXYLAMINE SUCCINATE (UNII: V9B19B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients	
Ingredient Name	**Strength**
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GELATIN (UNII: 2G86QN327L)	
POVIDONE (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
GLYCERIN (UNII: PDC6A3C0OX)	
SHELLAC (UNII: 46N107B71O)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	green	**Score**	no score
Shape	CAPSULE (CAPSULE SHAPED)	**Size**	24mm
Flavor		**Imprint Code**	72
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11673-979-08	8 in 1 BOX; Type 0: Not a Combination Product	07/13/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	07/13/2020	

Labeler - TARGET CORP (006961700)

Registrant - TIME CAP LABS INC (037052099)

Establishment

Name	Address	ID/FEI	Business Operations
MARKSANS PHARMA LTD		925822975	manufacture(11673-979)

Revised: 7/2020 TARGET CORP