# 8HR PAIN RELIEF- acetaminophen tablet, extended release TARGET CORPORATION

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### 704R Target Acetaminophen Extended-Release Tablets USP, 650 mg

#### Active ingredient (in each caplet)

Acetaminophen 650 mg

#### Purpose

Pain reliever/fever reducer

#### Uses

temporarily relieves minor aches and pains due to:

- muscular aches
- backache
- minor pain of arthritis
- toothache
- premenstrual and menstrual cramps
- headache
- the common cold

temporarily reduces fever

#### Warnings

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

- more than 6 caplets in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day 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 allergic to acetaminophen or any of the inactive ingredients in this product.

### Ask a doctor before use if you have liver disease.

Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin.

### Stop using and ask a doctor if

- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- new symptoms occur
- redness or swelling is present

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. (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.

# Directions

## Do not take more than directed. See overdose warning

adults and children 12 years and over:

- take 2 caplets every 8 hours with water
- swallow whole; do not crush, chew, split or dissolve
- do not take more than 6 caplets in 24 hours
- do not use for more than 10 days unless directed by a doctor

children under 12 years: do not use

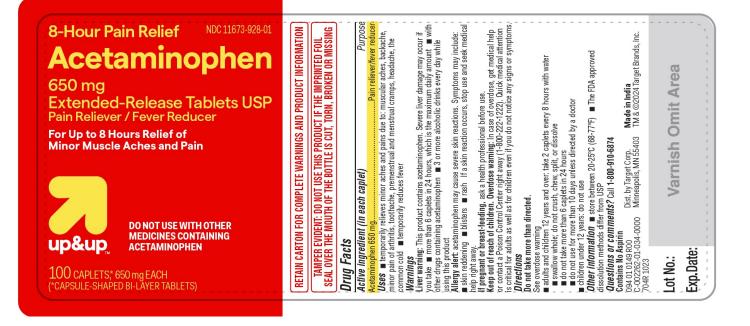
## Other information

- store between 20-25°C (68-77°F)
- The FDA approved Dissolution methods differ from USP

*Inactive ingredients* carnauba wax, hydroxyethyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, povidone, pregelatinized starch, sodium starch glycolate, titanium dioxide, triacetin

## **Questions or comments?**

Call **1-800-910-6874** 





8HR PAIN RELIEF acetaminophen tablet, extended release										
Product Information										
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:11673-928						
Route of Administration	ORAL									
Active Ingredient/Active Moiety										
Ingr		Basis of St	trength	Strength						

In	active Ingre	edients									
Ingredient Name											
sт	ARCH, CORN (U	NII: 08232NY3SJ)									
POVIDONE K30 (UNII: U725QWY32X)											
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)											
TRIACETIN (UNII: XHX3C3X673)											
CA	RNAUBA WAX (	UNII: R12CBM0EIZ)									
Τľ	TANIUM DIOXID	E (UNII: 15FIX9V2JP)									
МІ	CROCRYSTALLI	NE CELLULOSE (UNII: OP1R32D61U)									
SC	DIUM STARCH	GLYCOLATE TYPE A (UNII: H8AV0SQX4D)									
H١	DROXYETHYL C	ELLULOSE (140 MPA.S AT 5%) (UNII: 8136	Y38GY5	)							
M	AGNESIUM STE	RATE (UNII: 70097M6I30)									
P	roduct Chara	acteristics									
Co	olor	white (White to off white)		Score	no score						
Shape CAPSULE (Capsule-shaped tablet)				Size	19mm						
Flavor				Imprint Code	71						
Contains											
Pa	ackaging										
#	ltem Code	Package Description	ſ	Marketing Start Date	Marketing End Date						
1	NDC:11673-928- 01	1 in 1 CARTON	11/	30/2022							
1		100 in 1 BOTTLE; Type 0: Not a Combinatio Product	n								
Μ	arketing	Information									
	Marketing Category	Application Number or Monogra Citation	ph	Marketing Start Date	Marketing End Date						

Labeler - TARGET CORPORATION (006961700)

**Registrant -** TIME CAP LABORATORIES, INC. (037052099)

Establishment									
Name	Address	ID/FEI	<b>Business Operations</b>						
MARKSANS PHARMA LIMITED		925822975	manufacture(11673-928)						