# ACETAMINOPHEN- acetaminophen tablet, coated TARGET CORPORATION

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.

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#### 341R-TGT APAP 500 MG 11673-838

Active Ingredient (in each tablet)

acetaminophen 500 mg

hypromellose, mineral oil, povidone, pregelatinized starch, sodium starch glycolate\* stearic acid, titanium dioxide

\*may contain this ingredient

#### PAIN RELIEVER - FEVER REDUCER

Uses: temporarily relieves minor aches and pains due to: the common cold, headache, backache, minor pain of arthritis, toothache, muscular aches, premenstrual and menstrual cramps, temporarily reduces fever.

Directions: Do not take more than directed (see overdose warning)

adults and children 12 years and over: take 2 caplets every 4 to 6 hours while symptoms last

do not take more than 8 caplets in 24 hours

do not take for more than 10 days unless directed by a doctor

children under 12 years: do not use this adult extra strength product in children under 12 years of age; this will provide more than the recommended does (overdose) of acetaminophen any may cause lier damage.

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

more than 4,000 mg of acetaminophen in 24 hours

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.

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). Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.



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### **ACETAMINOPHEN**

acetaminophen tablet, coated

Product	Information
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Product TypeHUMAN OTC DRUGItem Code (Source)NDC:11673-838

ORAL

## **Active Ingredient/Active Moiety**

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Ingredient Name	Basis of Strength	Strength
ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D)	ACETAMINOPHEN	500 mg

Inactive Ingredients		
Ingredient Name	Strength	
MINERAL OIL (UNII: T5L8T28FGP)		
STARCH, CORN (UNII: O8232NY3SJ)		
HYPROMELLOSES (UNII: 3NXW29V3WO)		
PO VIDO NE, UNSPECIFIED (UNII: FZ989GH94E)		
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)		

Product Characteristics			
Color	white	Score	no score
Shape	OVAL	Size	17mm
Flavor		Imprint Code	TCL341
Contains			

I	Packaging			
#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:11673-838-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	0 1/0 1/20 20	
2	NDC:11673-838-05	50 in 1 BOTTLE; Type 0: Not a Combination Product	0 1/0 1/20 20	
3	NDC:11673-838-50	500 in 1 BOTTLE; Type 0: Not a Combination Product	0 1/0 1/20 20	
4	NDC:11673-838-26	225 in 1 BOTTLE; Type 0: Not a Combination Product	0 1/0 1/20 20	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	0 1/0 1/20 20	

# Labeler - TARGET CORPORATION (006961700)

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

Establishment				
Name	Address	ID/FEI	Business Operations	
TIME CAP LABORATORIES, INC.		037052099	manufacture(11673-838)	

Revised: 9/2019 TARGET CORPORATION