

**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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**340R-TARGET - APAP 325 MG 11673-841**

**ACTIVE INGREDIENT (IN EACH TABLET)**

Acetaminophen 325mg

povidone, pregelatinized starch, sodium starch glycolate\*, stearic acid

\*may contain this ingredient

**Purpose**

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

**DOSAGE & ADMINISTRATION**

**Directions**

do not take more than directed (see overdose warning)

adults and children 12 years and over

take 2 tablets (1,000 mg) every 6 hours while symptoms last

do not take more than 6 tablets (3,000 mg) in 24 hours, unless directed by a doctor

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 dose (overdose) of acetaminophen and may cause liver damage

Keep out of reach of children.

In case of accidental overdose, get medical help or contact a Poison Control Center. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

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

- more than 12 tablets 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.



acetaminophen tablet, coated

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11673-841
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg

### Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
POVIDONE (UNII: FZ989GH94E)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
STARCH, CORN (UNII: O8232NY3SJ)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	

### Product Characteristics

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	10 mm
<b>Flavor</b>		<b>Imprint Code</b>	TCL340
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-841-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	

### Marketing Information

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC monograph not final	part343	01/01/2020	

**Labeler** - TARGET CORPORATION (006961700)

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

### Establishment

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
TIME CAP LABORATORIES, INC.		037052099	manufacture(11673-841)

