

**EXTRA STRENGTH ACETAMINOPHEN- acetaminophen tablet  
TIME CAP LABORATORIES, INC.**

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**673T Timely Extra Strength Acetaminophen Gelcaps**

**Drug Facts**

**Active ingredient (in each gelcap)**

Acetaminophen USP 500 mg

**Purpose**

Pain reliever/fever reducer

**Uses**

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

**Warnings**

**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.

**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 use 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 gelcaps every 6 hours while symptoms last
- do not take more than 6 gelcaps in 24 hours, unless directed by a doctor
- do not use for more than 10 days unless directed by a doctor

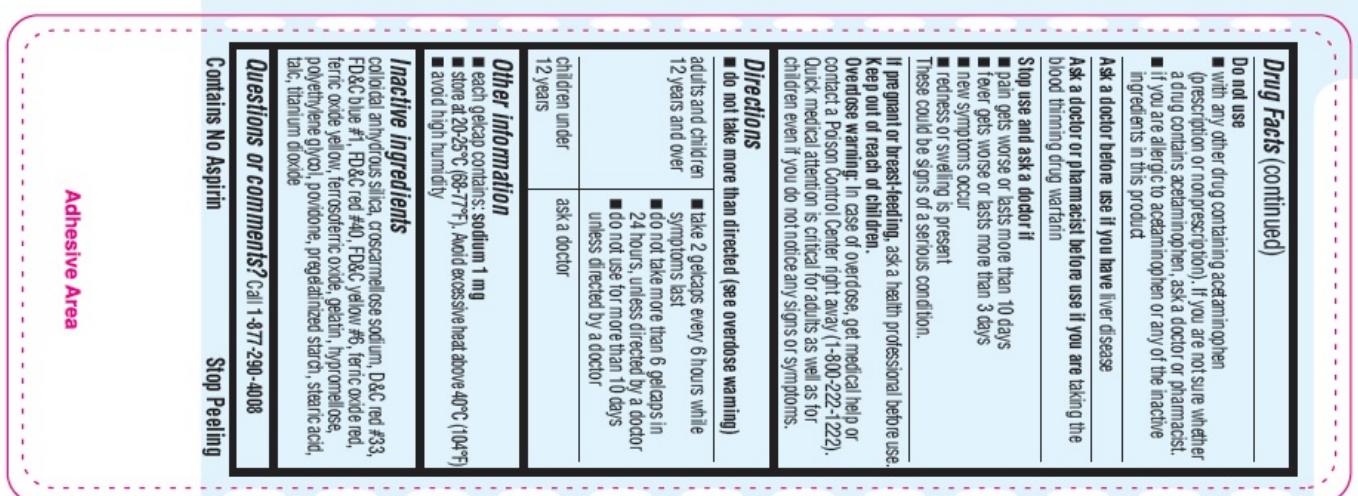
children under 12 years      ask a doctor

**Other information**

- each gelcap contains: **sodium 1 mg**
- store between 20-25°C (68-77°F). Avoid excessive heat above 40°C (104°F)
- avoid high humidity

**Inactive ingredients** colloidal anhydrous silica, croscarmellose sodium, D&C red #33, FD&C blue #1, FD&C red #40, FD&C yellow #6, ferric oxide red, ferric oxide yellow, ferrosferric oxide, gelatin, hypromellose, polyethylene glycol, povidone, pregelatinized starch, stearic acid, talc, titanium dioxide

**Questions or comments?** Call **1-877-290-4008**



# **EXTRA STRENGTH ACETAMINOPHEN**

## acetaminophen tablet

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDc:83934-673
<b>Route of Administration</b>	ORAL		

## **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

## Inactive Ingredients

Ingredient Name	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POVIDONE (UNII: FZ989GH94E)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

**FD&C RED NO. 40** (UNII: WZB9127XOA)

**FERRIC OXIDE YELLOW** (UNII: EX43802MRT)

**POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WJQ0SDW1A)

**STEARIC ACID** (UNII: 4ELV7Z65AP)

**TALC** (UNII: 7SEV7J4R1U)

**SILICON DIOXIDE** (UNII: ETJ7Z6XBU4)

**D&C RED NO. 33** (UNII: 9DBA0SBB0L)

**FD&C BLUE NO. 1** (UNII: H3R47K3TBD)

**FERRIC OXIDE RED** (UNII: 1K09F3G675)

**FERROSOFERRIC OXIDE** (UNII: XM0M87F357)

**GELATIN** (UNII: 2G86QN327L)

**CROSCARMELLOSE SODIUM** (UNII: M28OL1HH48)

**STARCH, CORN** (UNII: O8232NY3SJ)

**HYDROXYPROPYL CELLULOSE, UNSPECIFIED** (UNII: 3NXW29V3WO)

## Product Characteristics

<b>Color</b>	red (Red opaque and Blue gray opaque hard gelatin shells)	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	19mm
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83934-673-73	100 in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2026	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	02/01/2026	

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

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

## Establishment

Name	Address	ID/FEI	Business Operations
MARKSANS PHARMA LIMITED		677604129	manufacture(83934-673)

Revised: 1/2026

TIME CAP LABORATORIES, INC.