

**PAIN RELIEVER- acetaminophen tablet, film coated  
WALGREENS**

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**Walgreens 679R ACETAMINOPHEN TABLETS USP 500 MG-Easy to swallow red caplets**

**Drug Facts**

**Active ingredient (in each caplet)**

Acetaminophen 500 mg

**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

**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 caplets every 6 hours while symptoms last  
do not take more than 6 caplets in 24 hours, unless directed by a doctor

children under 12 years

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

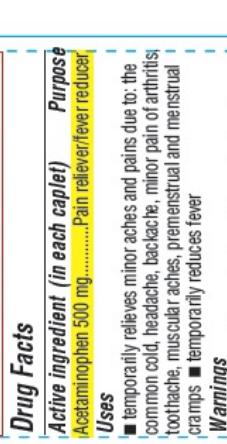
ask a doctor

**Other information**

store between 20-25°C (68-77°F)

**Inactive ingredients** carnauba wax, FD&C red no. 40 al. lake, FD&C yellow no. 6 al. lake, guar gum, hypromellose, light mineral oil, maltodextrin, povidone, pregelatinized starch, sodium starch glycolate, stearic acid, sucralose, talc, titanium dioxide

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

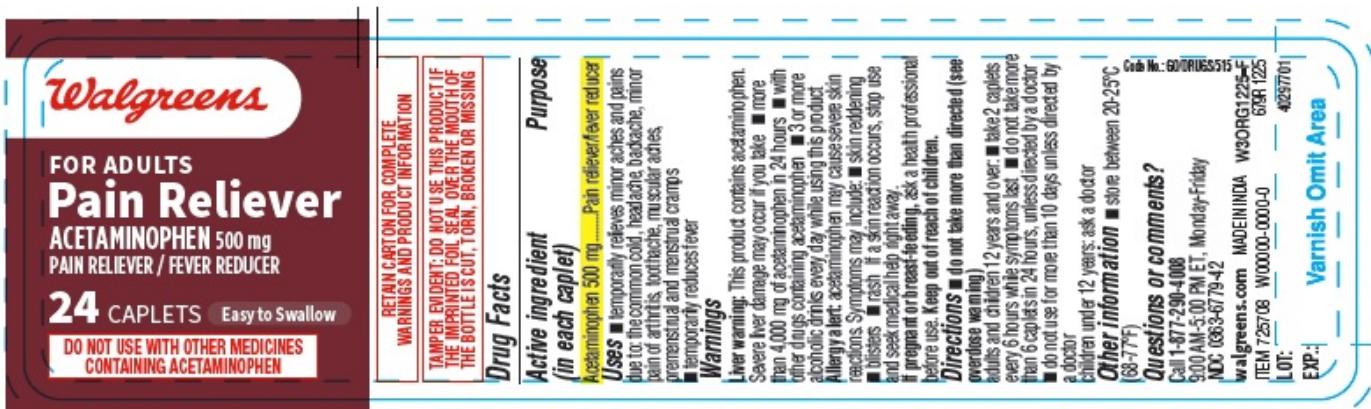


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Questions or comments? Call 1-877-290-4008  
9:00 AM-5:00 PM ET, Monday-Friday  
Walgreens.com  
NDC 0854-6779-01  
402761 Cap No. 500BLS55  
ITEM 725681  
W20RG1225-F  
679R1225  
LOT: Vanish On/Off Area

## Varnish/Print Omit Area





## PAIN RELIEVER

acetaminophen tablet, film coated

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-6779
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	500 mg

### Inactive Ingredients

Ingredient Name	Strength
FD&C YELLOW NO. 6 ALUMINUM LAKE (UNII: GYP6Z2JR6Q)	

GUAR GUM (UNII: E89I1637KE)
TALC (UNII: 7SEV7J4R1U)
MALTODEXTRIN (UNII: 7CVR7L4A2D)
STEARIC ACID (UNII: 4ELV7Z65AP)
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)
STARCH, CORN (UNII: 08232NY3SJ)
SUCRALOSE (UNII: 96K6UQ3ZD4)
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)
POVIDONE (UNII: FZ989GH94E)
FD&C RED NO. 40 ALUMINUM LAKE (UNII: 6T47AS764T)
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)
LIGHT MINERAL OIL (UNII: N6K5787QVP)
CARNAUBA WAX (UNII: R12CBM0EIZ)

## Product Characteristics

Color	red	Score	no score
Shape	OVAL (Capsule shaped tablets)	Size	18mm
Flavor		Imprint Code	TCL341
Contains			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-6779-42	1 in 1 CARTON	02/01/2026	
1		24 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:0363-6779-01	1 in 1 CARTON	02/01/2026	
2		100 in 1 BOTTLE; Type 0: Not a Combination Product		

## Marketing Information

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

**Labeler** - WALGREENS (008965063)

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

## Establishment

Name	Address	ID/FEI	Business Operations
MARKSANS PHARMA LIMITED		925822975	manufacture(0363-6779)

