REGULAR STRENGTH ACETAMINOPHEN- acetaminophen tablet CARDINAL HEALTH

340R CARDINAL HEALTH 70000 0092 Acetaminophen Tablets

DRUG FACTS

Active ingredient (in each tablet)

Acetaminophen 325 mg

Purpose

Pain reliever/fever reducer

Uses

temporarily relieves minor aches and pains due to:

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

temporarily reduces fever

Warnings

Liver warning: This product contains acetaminophen. The maximum daily dose of this product is 10 tablets (3,250 mg) in 24 hours for adults or 5 tablets (1,625 mg) in 24 hours for children. Severe liver damage may occur if

- adult takes more than 4,000 mg of acetaminophen in 24 hours
- child takes more than 5 doses in 24 hours, which is the maximum daily amount
- taken with other drugs containing acetaminophen
- adult has 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 in adults
- pain gets worse or lasts more than 5 days in children under 12 years
- 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.

In case of accidental overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Prompt 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

adults and children 12 years and over

- take 2 tablets (650 mg) every 4 to 6 hours while symptoms last
- do not take more than 10 tablets (3,250 mg) in 24 hours, unless directed by a doctor
- do not use for more than 10 days unless directed by a doctor

children 6 years to under 12 years

- take 1 tablet (325 mg) every 4 to 6 hours while symptoms last
- do not take more than 5 tablets (1,625 mg) in 24 hours
- do not use for more than 5 days unless directed by a doctor

children under 6 years ask a doctor

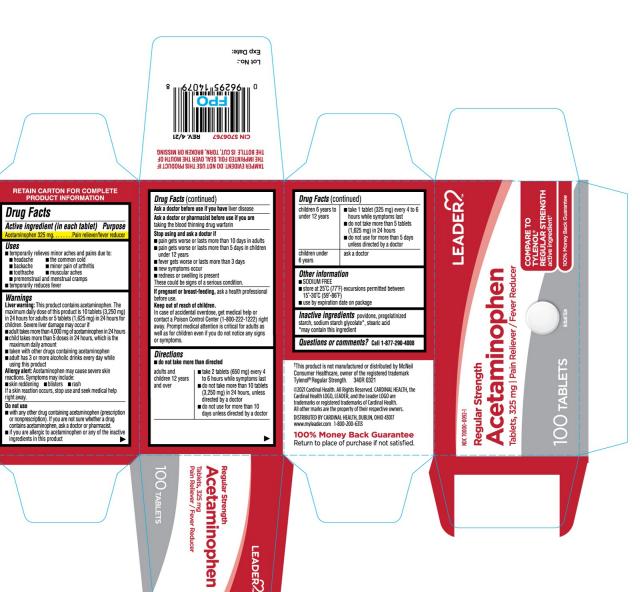
OTHER INFORMATION

Other information

- SODIUM FREE
- store at 25°C (77°F) excursions permitted between 15°-30°C (59°-86°F)
- use by expiration date on package

Inactive ingredients povidone, pregelatinized starch, sodium starch glycolate*, stearic acid *may contain this ingredient

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



Drug Facts

Uses

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In hadache

I the common cold

Backche

I minor pain of arthritis

I premenstrual and menistrual cramps

I temporarily reduces fever

Uses

right away.



REGULAR STRENGTH ACETAMINOPHEN

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70000-0092
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg	

Inactive Ingredients				
Ingredient Name	Strength			
STARCH, CORN (UNII: 08232NY3SJ)				
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)				
POVIDONE (UNII: FZ989GH94E)				
STEARIC ACID (UNII: 4ELV7Z65AP)				

Product Characteristics			
Color	white	Score	2 pieces
Shape	ROUND	Size	10mm
Flavor		Imprint Code	TCL;340
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70000- 0092-1	1 in 1 CARTON	05/05/2021	
1		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	05/05/2021	

Labeler - CARDINAL HEALTH (063997360)

Registrant - TIME CAP LABORATORIES, INC. (037052099)

Establishment				
Name	Address	ID/FEI	Business Operations	
TIME CAP LABORATORIES, INC.		037052099	manufacture(70000-0092)	

Revised: 1/2024 CARDINAL HEALTH