ACETAMINOPHEN- acetaminophen tablet extended release tablet, extended release MUSCLE ACHES AND PAINS ACETAMINOPHEN EXTENDED RELEASEacetaminophen tablet, extended release TIME CAP LABORATORIES, INC.

Timely 699R 704R Acetaminophen Extended-Release Tablets USP, 650 mg

Active ingredient (in each caplet)

Acetaminophen 650 mg

Purpose

Pain reliever/fever reducer

Uses

For Arthritis Pain

temporarily relieves minor aches and pains due to:

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

temporarily reduces fever

For Muscle Aches & Pains

temporarily relieves minor aches and pains due to:

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

temporarily reduces fever

Warnings

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

- more than 6 caplets in 24 hours, which is the maximum daily amount
- 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 using 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

For Arthritis Pain

Do not take more than directed. See overdose warning

adults:

- take 2 caplets every 8 hours with water
- swallow whole; do not crush, chew, split or dissolve
- do not take more than 6 caplets in 24 hours
- do not use for more than 10 days unless directed by a doctor

under 18 years of age: ask a doctor

For Muscle Aches & Pains

Do not take more than directed. See overdose warning

adults and children 12 years and over:

- take 2 caplets every 8 hours with water
- swallow whole; do not crush, chew, split or dissolve

- do not take more than 6 caplets in 24 hours
- do not use for more than 10 days unless directed by a doctor

children under 12 years: do not use

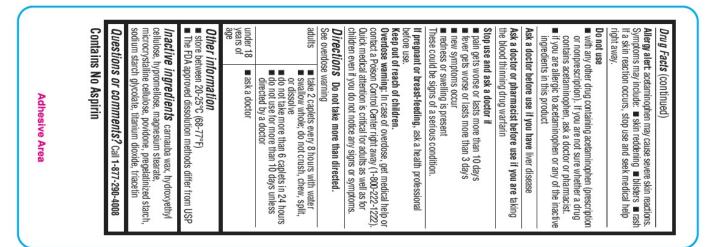
Other information

- store between 20-25°C (68-77°F)
- The FDA approved Dissolution methods differ from USP

Inactive ingredients carnauba wax, hydroxyethyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, povidone, pregelatinized starch, sodium starch glycolate, titanium dioxide, triacetin

Questions or comments?

Call **1-877-290-4008**



Inside



*Compare to T	Ulenol ^{® Ske} Drug Facts	Drug Facts (continued)	Drug Facts (continued)		
	ingredient Active ingredient (in each caplet) Purpu Acetaminophen 650 mgPain reliever/tever redu Uses I temporarily relieves minor aches and pains due to:		swallow whole; do not crush, chew, split, or dissolve do not take more than 6 caplets in 24 hours do not take more than 10 days unless directed by a doctor under 18 aks 4 adoctor		
Arthritis Pain Relief	 minor pain of arthritis muscular aches backache premenstrual and menstrual cramp the common cold headache toothache 		years of age		
Acetaminophen	temporarily reduces fever Warnings	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	Other information store between 20-25°C (68-77°F) The FDA approved dissolution methods differ from USP		
Extended-Release Tablets USP, 650 i P <mark>ain Reliever/Fever Reducer</mark>	more than 6 caplets in 24 hours, which is the maxim daily amount	These could be signs of a serious condition.	Inactive ingredients carnauba wax, hydroxyethyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, povidone, pregelatinized starch, sodium starch glycolate, titanium dioxide, triacetin Auestione or comments? Cell 1.st7-200-4008		
Lasts up to 8 hours	 with other drugs containing acetaminophen 3 or more alcoholic drinks every day while using this product Allergy alert: acetaminophen may cause severe skin 	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			
minor arthritis pain Actual s	size reactions. Symptoms may include: ■ skin reddening ■ blisters ■ rash	contact a Poison Control Center right away (1-800-222-1222). Quick medical attention is critical for adults as well as for	This product is not manufactured or distributed by Johnson & Johnson		
DO NOT USE WITH OTHER MEDICINES CONTAINING ACETAMINOPHEN		Directions Do not take more than directed. See overdose warning	Consumer Inc., McNeil Consumer Healthcare division, owner of the registered trademark Tylenol [®] & Arthritis Pain Distributed by: Time-Cap Labs, Inc.		
*Capsule-Shaped Bi-Layer Tablets 650 mg I	with any other drug containing acetaminophen	► adults ■ take 2 caplets every 8 hours with water ►	7 Michael Avenue, Farmingdale, NY 11735 699R 0723 Made in India	Lot No.:	

NDC 49483-704-05	CGR COMPLETE IRMATION IRMATION A THE MOUTH OF THE KEN OR MISSING THE MOUTH OF THE KEN OR MISSING FUE contains acetaminophen. Table of arthritis sand pains due to: "val cramps • headache es fever te contains acetaminophen. "a nore than 6 caplets in nount = with other drugs alcoholic drinks every day severe skin reactions. of = blisters = rash et mad 3 days = new is present. These could be the professional before use. the professional before use. the professional before use. the professional before use. admission Control Center right petes in 24 hours = det 2 caplets of by a doctor children of by a doctor children of by a doctor children to the farmingdale, Avenue, farmingdale,
Muscle Aches & Pain Acetaminophen Extended-Release Tablets USP, 650 mg Pain Reliever/Fever Reducer For up to 8 hours relief of Minor Muscle Aches & Pain	PC CARTON FOR THI DO NOT USE THI SE ANDD INFORMAN SEAL OUT USE THI SEAL OUT USE THI SEAL OUT USE THI SEAL OUT AND SEAL OUT AND SEAL OUT AND SEAL OUT AND SEAL OUT AND SEAL OUT AND ADDREND AND ADDR
DO NOT USE WITH OTHER MEDICINES CONTAINING ACETAMINOPHEN 50 Caplets* Capsule-Shaped Bi-Layer Tablets 650 mg EACH	READ AND KEE READ AND KEE MARNING TAMPER EVIDENT: THE IMPRINTED FOIL BOTTLE IS CUT, BOTTLE IS CU



Contains	Inactive ing carnauba wax, h magnesium stea povidone, pregel titanium dioxide, duestions o	Other info ■ store betwe ■ The FDA ap USP	under 18 years of age	adults	Directions Do not take m See overdose	If pregnant or breast-feed before use. Overdose warning: In cass or contact a Poisson Contro right away. Quick medical well as for children even if symptoms.	■ new symptoms occurs redness or swelling is These could be signs of	Stop using an ■ pain gets w ■ fever gets v	Ask a doctor the blood thir	or nonpress contains ac if you are a inactive ing Ask a doctor	Drug Facts
lo Aspirin Adhesive Area	Inactive ingredients carnauba wax, hydroxyethyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, povidone, pregelatinized starch, sodium starch glycolate, titanium dioxide, triacetin Questions or comments? Call 1-877-290-4008	<i>r information</i> e between 20-25°C (68-77°F) FDA approved Dissolution methods differ from	■ ask a doctor	 take 2 caplets every 8 hours with water swallow whole; do not crush, chew, split or dissolve do not take more than 6 caplets in 24 hours do not use for more than 10 days unless directed by a doctor 	7S more than directed. se warning	If pregnant or breast-feeding, ask a health professional before use. Neep out of reach of children. Overdose warning: In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.	swelling is present swelling is present signs of a serious condition.	Stop using and ask a doctor if ■ pain gets worse or lasts more than 10 days ■ fever gets worse or lasts more than 3 days	Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin	or nonprescription), it 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 before use if you have liver disease	









ACETAMINO								
acetaminophen ta	ablet extend	ed release tablet, exte	nded release					
Product Inform	mation							
Product Type		HUMAN OTC DRUG	Item Code (So	urce)	NDC:494	83-699		
Route of Adminis	stration	ORAL						
Active Ingredie	ent/Active	Moiety						
	Ingre	edient Name		Basis of S	Strength	Strength		
ACETAMINOPHEN (UNII: 36209ITL	9D) (ACETAMINOPHEN - UNI	I:362O9ITL9D)	ACETAMINOP	HEN	650 mg		
Inactive Ingree	dients							
		Ingredient Name				Strength		
STARCH, CORN (UN	-							
POVIDONE K30 (UN								
MAGNESIUM STEARATE (UNII: 70097M6I30) HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)								
TRIACETIN (UNII: XH		UIVII: SIVAVZ9V3VVU)						
CARNAUBA WAX (U		17)						
		E (UNII: OP1R32D61U)						
		PE A (UNII: H8AV0SQX4D)						
HYDROXYETHYL CI	ELLULOSE (14	0 MPA.S AT 5%) (UNII: 81	36Y38GY5)					
Product Chara	ctoristics							
		to off white)	C c c m c		20.0	coro		
Color	white (White CAPSULE	to off white)	Score Size		10 s	core		
Shape	CAPSULE			Cada		111		
Flavor			Imprint	Lode	71			
Contains								
Packaging								
# Item Code	Pac	kage Description		ting Start Date		ting End ate		
	400 in 1 BOTT Product	LE; Type 0: Not a Combinat	tion 10/04/2021	L				
	100 in 1 BOTT Product	LE; Type 0: Not a Combinat	tion 10/04/2023					
05	50 in 1 CARTO		10/04/2023	1				
3	Product	Type 0: Not a Combination						
4	225 in 1 BOTT Product	LE; Type 0: Not a Combinat	tion 11/07/2022	2				
26 NDC:49483-699-	Product							

MUSCLE ACHES AND PAINS ACETAMINOPHEN EXTENDED RELEASE acetaminophen tablet, extended release Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:49483-704 Route of Administration ORAL V V Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength

Ingredient Name	Basis of Strength	Streng
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	650 mg

Ingredient Name	Strength
STARCH, CORN (UNII: 08232NY3SJ)	Stiength
POVIDONE K30 (UNII: U725QWY32X)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
TRIACETIN (UNII: XHX3C3X673)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	
HYDROXYETHYL CELLULOSE (140 MPA.S AT 5%) (UNII: 8136Y38GY5)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
Product Characteristics	

Color	white (White to off white)	Score	no score
Shape	CAPSULE (Capsule-shaped tablet)	Size	19mm
Flavor		Imprint Code	71
Contains			

Packaging # Item Code Package Description Marketing Start Date Marketing End Date

1 05	1 in 1 CARTON	11/07/2022				
1	0 in 1 BOTTLE; Type 0: Not a Combination roduct					
Marketing Information						
Marketing	Information					
Marketing Marketing Category	Information Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
Marketing	Application Number or Monograph	-	-			

Labeler - TIME CAP LABORATORIES, INC. (037052099)

Registrant - TIME CAP LABORATORIES, INC. (037052099)

Establishment							
Name	Address	ID/FEI	Business Operations				
MARKSANS PHARMA LIMITED		925822975	manufacture(49483-699, 49483-704)				

Revised: 3/2024

TIME CAP LABORATORIES, INC.