ARTHRITIS PAIN ACETAMINOPHEN- acetaminophen tablet
TARGET CORPORATION

699R Target 11673 673 Acetaminophen Extended Release Tablets USP, 650 mg

DRUG FACTS

Active ingredient (in each caplet)

Acetaminophen 650 mg

Purpose

Pain reliever/fever reducer

Uses

temporarily relieves minor aches and pains due to:

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

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 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	• 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

Other information

- store between 20-25ºC (68-77ºF)
- 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**

OR

Call **1-800-910-6874**

Drug Facts (continued)

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
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Questions or comments? Call 1-800-910-6874

Contains No Aspirin

Adhesive Area

Compare to active ingredient in Tylenol® 8HR†

Arthritis Pain Acetaminophen

650 mg Extended-Release Tablets USP
Pain Reliever / Fever Reducer

- For the temporary relief of minor arthritis pain
- Lasts up to 8 hours

PACKAGE NOT CHILD RESISTANT

Actual Size

up&up™

100 CAPLETS* 650 mg EACH
(*CAPSULE-SHAPED BI-LAYER TABLETS)

NDC 11673-673-01

DO NOT USE WITH OTHER MEDICINES CONTAINING ACETAMINOPHEN

TAMPER EVIDENT: DO NOT USE THIS PRODUCT IF THE IMPRINTED FOIL SEAL OVER THE MOUTH OF THE BOTTLE IS CUT, TORN, BROKEN OR MISSING

Drug Facts

Active ingredient (in each caplet)	Purpose
Acetaminophen 650 mg	Pain reliever/fever reducer

Uses
■ 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

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- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Drug Facts (continued under label)

This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., McNeil Consumer Healthcare division, owner of the registered trademark Tylenol® 8HR
Arthritis Pain
094 01 0496 R00 C-002262-01-034-0000
Distributed by Target Corporation
Minneapolis, MN 55403
Made in India
TM & ©2024 Target Brands, Inc.
699R 1023

3 11673 67301 0

FEEL HERE

Lot No.:

Exp.Date:

Varnish Omit Area

RETAIN CARTON FOR COMPLETE PRODUCT INFORMATION

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

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Other information ■ **TAMPER EVIDENT: DO NOT USE THIS PRODUCT IF THE IMPRINTED FOIL SEAL OVER THE MOUTH OF THE BOTTLE IS CUT, TORN, BROKEN OR MISSING**

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

Contains no aspirin

094 01 3533 R00 C-001744-01-039 This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., owner of the registered trademark Tylenol®.
Dist. by Target Corp., Mpls., MN 55403
Made in India code No.: 60/DRUGS/515
TM & ©2022 Target Brands, Inc.
699R 0921
9BL442
Lot No.:
Exp. Date:

Compare to active ingredient in **Tylenol® 8HR Arthritis Pain†**

arthritis pain
acetaminophen

extended-release tablets USP, 650 mg
pain reliever/fever reducer

for the temporary relief of minor arthritis pain
lasts up to 8 hours

up&up™

225 CAPLETS
CAPSULE-SHAPED BI-LAYER TABLETS

NDC 11673-673-26

225 CAPLETS

ARTHRITIS PAIN ACETAMINOPHEN

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-673
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg

Inactive Ingredients

Ingredient Name	Strength
POVIDONE K30 (UNII: U725QWY32X)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
HYDROXYETHYL CELLULOSE (140 CPS AT 5%) (UNII: 8136Y38GY5)	
STARCH, CORN (UNII: O8232NY3SJ)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
TRIACETIN (UNII: XHX3C3X673)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-673-26	1 in 1 CARTON	12/17/2021	
1		225 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:11673-673-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	11/30/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA215486	12/17/2021	

Labeler - TARGET CORPORATION (006961700)

Registrant - TIME CAP LABORATORIES, INC. (037052099)

Establishment

Name	Address	ID/FEI	Business Operations
MARKSANS PHARMA LIMITED		925822975	manufacture(11673-673)

Revised: 3/2024 TARGET CORPORATION