ACETAMINOPHEN- acetaminophen tablet, film coated
TARGET CORPORATION

342R TARGET - APAP 500 MG TABLETS - 11673-342

Drug Facts

Active ingredient (in each tablet)

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 tablets every 6 hours while symptoms last
- do not take more than 6 tablets 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

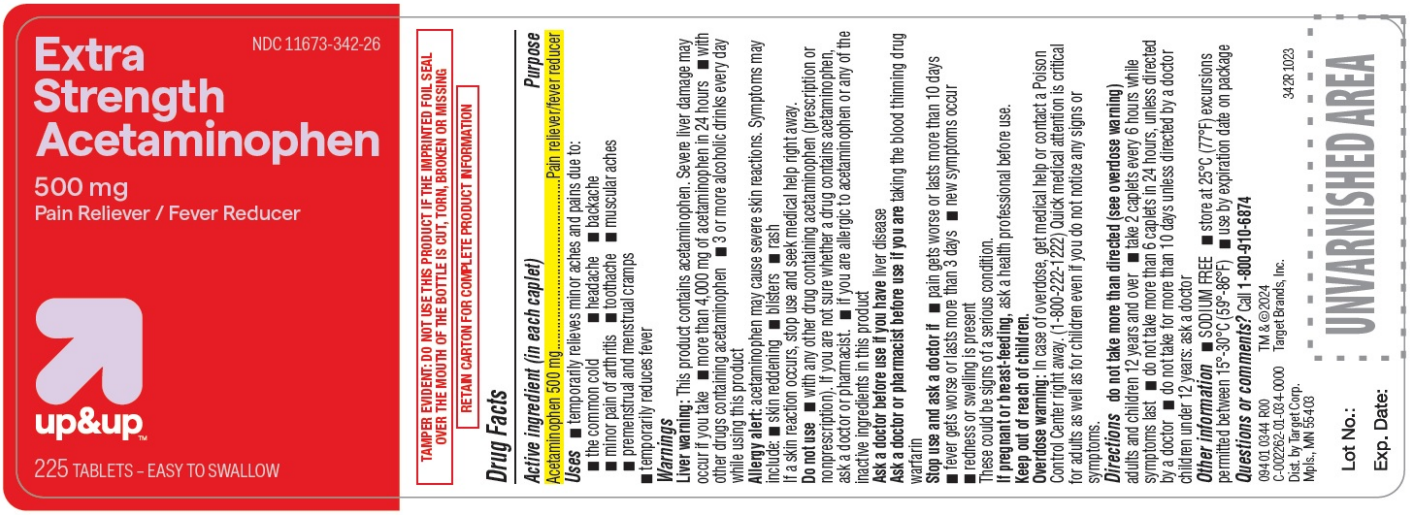
Other information

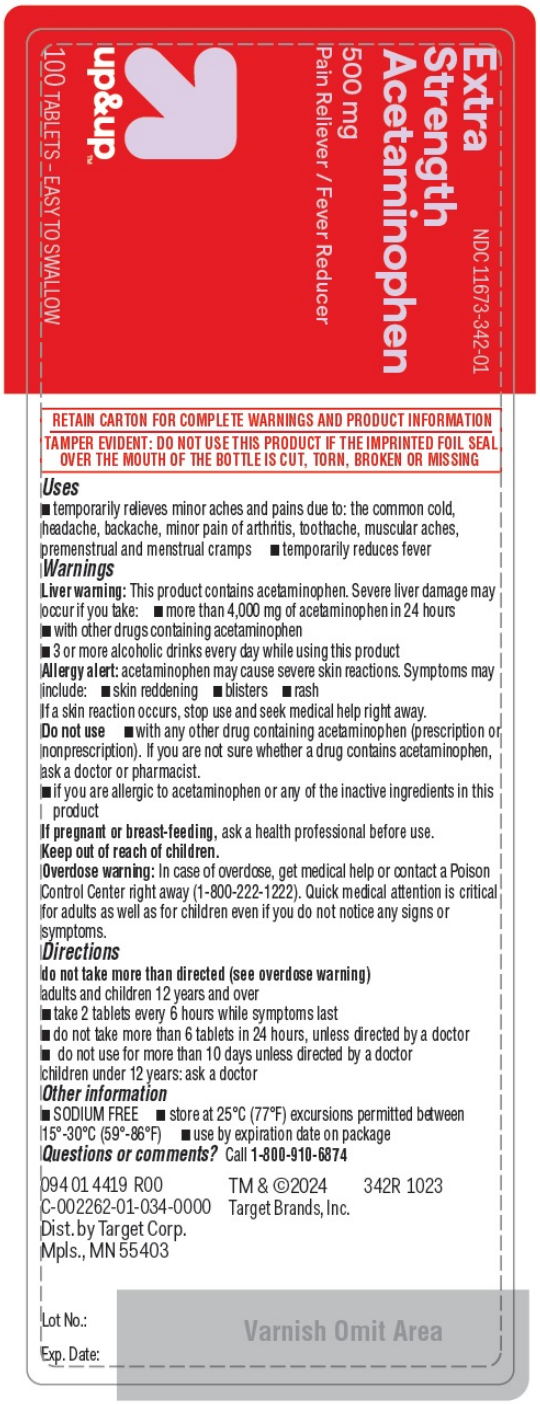
- SODIUM FREE
- store at 25°C (77°F) excursions permitted between 15°-30°C (59°-86°F)
- use by expiration date on package

Inactive ingredients carnauba wax, FD&C red #40 aluminum lake, hypromellose, polyethylene glycol, povidone, pregelatinized starch, sodium starch glycolate*, stearic acid, sucralose, titanium dioxide

*may contain this ingredient

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




Extra Strength Acetaminophen
500 mg
Pain Reliever / Fever Reducer
up&up
100 TABLETS - EASY TO SWALLOW
NDC 11673-342-01
RETAIN CARTON FOR COMPLETE WARNINGS AND PRODUCT 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
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.
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Questions or comments? Call 1-800-910-6874
094 01 4419 R00
C-002262-01-034-0000
Dist. by Target Corp.
Mpls., MN 55403
TM & ©2024
Target Brands, Inc.
342R 1023
Lot No.:
Exp. Date:
Varnish Omit Area


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Questions or comments?
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This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., Maker of Consumer Healthcare Division, owner of the registered trademark Extra Strength Tylenol®.
Distributed by Target Corporation
Minneapolis, MN 55403
TM & ©2024 Target Brands, Inc.
342R 1023
Extra Strength Acetaminophen
500 mg
Pain Reliever / Fever Reducer
up&up
Compare to active ingredient in Extra Strength Tylenol®†
NDC 11673-342-26
Extra Strength Acetaminophen
500 mg
Pain Reliever / Fever Reducer
225 TABLETS - EASY TO SWALLOW
Actual Size 225 Tablets
Coating Omit
Coating/Print Omit Area
Lot No.:
Exp.Date:


Lot No.: Coating/Print Omit Area
Exp. Date:
3 49483 54201 7
0940I 419 R00
C-00252-01-034-0000
Coating Omit
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342R 1023
Satisfaction guaranteed – Love it or your money back.
Distributed by Target Corporation
Minneapolis, MN 55403
TM & ©2024 Target Brands, Inc.
NDC 11673-342-01
Compare to active ingredient in Extra Strength Tylenol®
Extra Strength Acetaminophen
500 mg
Pain Reliever / Fever Reducer
Actual Size
100 Tablets
100 TABLETS – EASY TO SWALLOW
up&up
Coating Omit
Coating/Print Omit Area
Coating Omit

ACETAMINOPHEN

acetaminophen tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-342
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL 8000 (UNII: Q662QK8M3B)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
STARCH, CORN (UNII: O8232NY3SJ)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
POVIDONE (UNII: FZ989GH94E)	

Product Characteristics

Color	red	**Score**	no score
Shape	ROUND	**Size**	11mm
Flavor		**Imprint Code**	TCL342
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-342-42	24 in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2019	
2	NDC:11673-342-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2019	
3	NDC:11673-342-26	225 in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC Monograph Drug	M013	06/01/2019	

Labeler - TARGET CORPORATION (006961700)

Registrant - TIME CAP LABORATORIES INC (037052099)

Establishment

Name	Address	ID/FEI	Business Operations
TIME CAP LABORATORIES, INC		037052099	manufacture(11673-342)

Revised: 9/2024 TARGET CORPORATION