

**ACETAMINOPHEN- acetaminophen tablet, film coated**  
**TARGET CORPORATION**

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**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<sup>™</sup>

225 TABLETS - EASY TO SWALLOW

NDC 11673-342-26

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

**RETAIN CARTON FOR COMPLETE PRODUCT INFORMATION**

**Drug Facts**

**Active ingredient (in each caplet)**  
Acetaminophen 500 mg.....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 redness ■ 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 ■ do not take 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  
**Questions or comments?** Call **1-800-910-6874**

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Dist. by Target Corp.  
Mpls., MN 55403

342R 1023

**UNVARNISHED AREA**

Lot No.:  
Exp. Date:

# Extra Strength Acetaminophen

500 mg  
Pain Reliever / Fever Reducer



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

**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

**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  
**Questions or comments?** Call 1-800-910-8874

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 Mpls., MN 55403

Varnish Omit Area

Lot No.:  
Exp. Date:

**up&up**  
 500 mg  
**Extra Strength Acetaminophen**  
 Pain Reliever / Fever Reducer

This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., McKelown Corporate Healthcare Division, owner of the registered trademark Extra Strength Tylenol®.  
 Distributed by Target Corporation 342R.1023  
 Minneapolis, MN 55403  
 TM & ©2024 Target Brands, Inc.

**Questions or comments?**  
 Call 1-800-910-8874

**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, hydroxypropyl, polyethylene glycol, povidone, pregelatinized starch, sodium starch glycolate, stearic acid, sucralose, titanium dioxide may contain this ingredient

**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

**Drug Facts (continued)**  
**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.

**Drug Facts**  
**Active Ingredient (in each tablet) Purpose**  
 Acetaminophen 500 mg..... 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.

Lot No.:  
Exp. Date:  
Coating/Print Omit Area

094.01.4419.R00  
C-002262-01-034-0000

NDC 11673-342-01

Compare to active ingredient in Extra Strength Tylenol®  
**Extra Strength Acetaminophen**

**up&up**  
 500 mg  
 Pain Reliever / Fever Reducer  
 225 TABLETS - EASY TO SWALLOW

NDC 11673-342-01

Compare to active ingredient in Extra Strength Tylenol®  
**Extra Strength Acetaminophen**

**up&up**  
 500 mg  
 Pain Reliever / Fever Reducer  
 225 TABLETS - EASY TO SWALLOW

Coating Omit

Coating/Print Omit Area

Actual Size 225 Tablets

Actual Size 225 Tablets



Lot No. Coating/Print  
Exp. Date:



RETAIN CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

### Drug Facts

**Active ingredient (in each tablet)** **Purpose**  
Acetaminophen 500 mg...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
  - hives
  - itching
  - swelling of the face, lips, tongue, or throat
  - difficulty breathing
- 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

### Drug Facts (continued)

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

do not take more than directed (see overdose warning)

- |                                       |  |
|---------------------------------------|--|
| adults and children 12 years and over | <ul style="list-style-type: none"> <li>take 2 tablets every 6 hours while symptoms last</li> <li>do not take more than 6 tablets in 24 hours, unless directed by a doctor</li> <li>do not use for more than 10 days unless directed by a doctor</li> </ul> |
| children under 12 years               | ask a doctor   |

### Drug Facts (continued)

#### 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

camellia 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 and comments?

Call 1-800-910-6874

\*This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., McNeil Consumer Healthcare Division, owner of the registered trademark Extra Strength Tylenol®

342R 1023

Satisfaction guaranteed - Love It or your money back.

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Minneapolis, MN 55403  
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Coating Omit

NDC187 384201

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

Coating/Print Omit Area

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

Coating Omit

NDC 11673-342-42  
extra strength  
**acetaminophen**  
tablets, 500 mg  
pain reliever/fever reducer



24 TABLETS  
easy-to-swallow

**IMPORTANT INFORMATION:** DO NOT USE THIS PRODUCT IF THE LABEL IS NOT ON THE BOTTLE. OVERUSE CAN CAUSE LIVER DAMAGE.

**NEVER GIVE TO A CHILD WHO DOES NOT HAVE THE 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.

**Warnings** — This product contains acetaminophen. Do not use if you are allergic to acetaminophen or any of the other ingredients listed on the label.

**Directions** — Adults and children 12 years of age and older: — Take 1 or 2 tablets every 4 to 6 hours, as needed, but do not take more than 8 tablets in 24 hours, which is the maximum daily amount. — Do not use with other drugs containing acetaminophen in 3 or more alcoholic drinks every day while using this product.

**Other Warnings** — Acetaminophen may cause severe liver damage if you take more than 8 tablets in 24 hours. — Do not use with other drugs containing acetaminophen. — 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. — Do not use if you are allergic to acetaminophen or any of the other ingredients listed on the label. — Do not use for more than 10 days for pain unless directed by a doctor.

**Other Facts** (continued under label)  
064-01-6640-800 C-800874-01-073  
Upjohn, Inc., Kalamazoo, MI 49001  
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Questions? Call 1-800-810-5874

3428119  
**FEEL HERE** Lot No.  
**FOR MORE** Expiration Date  
**FACTS**

acetaminophen tablet, film coated

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11673-342
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

## Inactive Ingredients

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

## Product Characteristics

<b>Color</b>	red	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	11mm
<b>Flavor</b>		<b>Imprint Code</b>	TCL342
<b>Contains</b>			

## 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
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OTC Monograph Drug	M013	06/01/2019	
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**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