ACETAMINOPHEN- acetaminophen tablet
TARGET CORPORATION

Extra strength
Acetaminophen gelcaps, 500mg
Rapid release
Pain reliever/fever reducer
For Adults

Active ingredient (in each gelcap)

Acetaminophen USP, 500mg

Purposes

Pain reliever/fever reducer

Uses

- temporarily relieves minor aches and pains due to:
- headache
- muscular aches
- backache
- minor pain of arthritis
- the common cold
- toothache
- premenstrual and menstrual cramps
- temporarily reduces fever

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.

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

- store at 20° - 25°C (68° - 77°F). See USP Controlled Room Temperature
- avoid high humidity
- see end panel for expiration date and lot number

Inactive Ingredients

ammonium hydroxide, black iron oxide, colloidal silicon dioxide, croscarmellose sodium, D&C red #33, FD&C blue #1, FD&C red #40, FD&C yellow #6, gelatin, hydroxypropyl cellulose, hydroxypropyl methyl cellulose, hypromellose, iron oxide red, isopropyl alcohol,
n-butyl alcohol, polyethylene glycol, povidone, pregelatinized starch, propylene glycol, shellac glaze, stearic acid, titanium dioxide, yellow iron oxide.

Questions or Comments?

call **1-877-770-3183** Mon-Fri 8:00 AM EST to 5:00 PM PST.

Principal Display Panel

code #	:
size	:1+3/4 X 1+3/4 X 3+3/8
ref #	:PP180604B view
material	:.016 SBS date

COATING FREE AREA

700000003230

200000004938

Drug Facts (continued)

Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin.

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Drug Facts (continued)

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Satisfaction guaranteed – Love it or your money back.

NDC 11673-167-05

Distributed by Target Corporation
Minneapolis, MN 55403
Made in India

ASPIRIN FREE

2000000004938
700000003230

COATING FREE AREA

up&up

Extra Strength Acetaminophen

500 mg Gelcaps, Rapid Release Pain Reliever / Fever Reducer

For Adults

50 GELCAPS

Actual Size 50 Gelcaps

Compare to active ingredient in Extra Strength Tylenol® Rapid Release Gels*

094 01 0548 R00 C-002262-01-034-0000

8 13874 02064 3

Lot

Exp.

COATING FREE AREA

READ AND KEEP CARTON FOR COMPLETE WARNINGS AND INFORMATION

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Drug Facts

Active ingredient (in each gelcap)	Purpose
Acetaminophen USP, 500 mg	Pain reliever/fever reducer

Uses
- temporarily relieves minor aches and pains due to:
  - headache
  - muscular aches
  - backache
  - minor pain of arthritis
  - the common cold
  - toothache
  - premenstrual and menstrual cramps
- temporarily reduces fever

Warnings

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

Compare to active ingredient in Extra Strength Tylenol® Rapid Release Gels*

Extra Strength Acetaminophen

500 mg Gelcaps, Rapid Release Pain Reliever / Fever Reducer

For Adults

up&up™

100 GELCAPS

Actual Size
100 Gelcaps

Important: Read all product information before using.

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

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(CONTINUED ON BACK OF LABEL)

NDC 11673-167-10
Dist. by Target Corporation
Minneapolis, MN 55403
Made in India

0 13874 02065 8
094 01 0547 R00 C-002262-01-034-0000
200000004939
700000003231

LOT

EXP

PEEL HERE

ASPIRIN FREE

Inside (adhesive side)

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 use and ask a doctor if

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Questions or comments?

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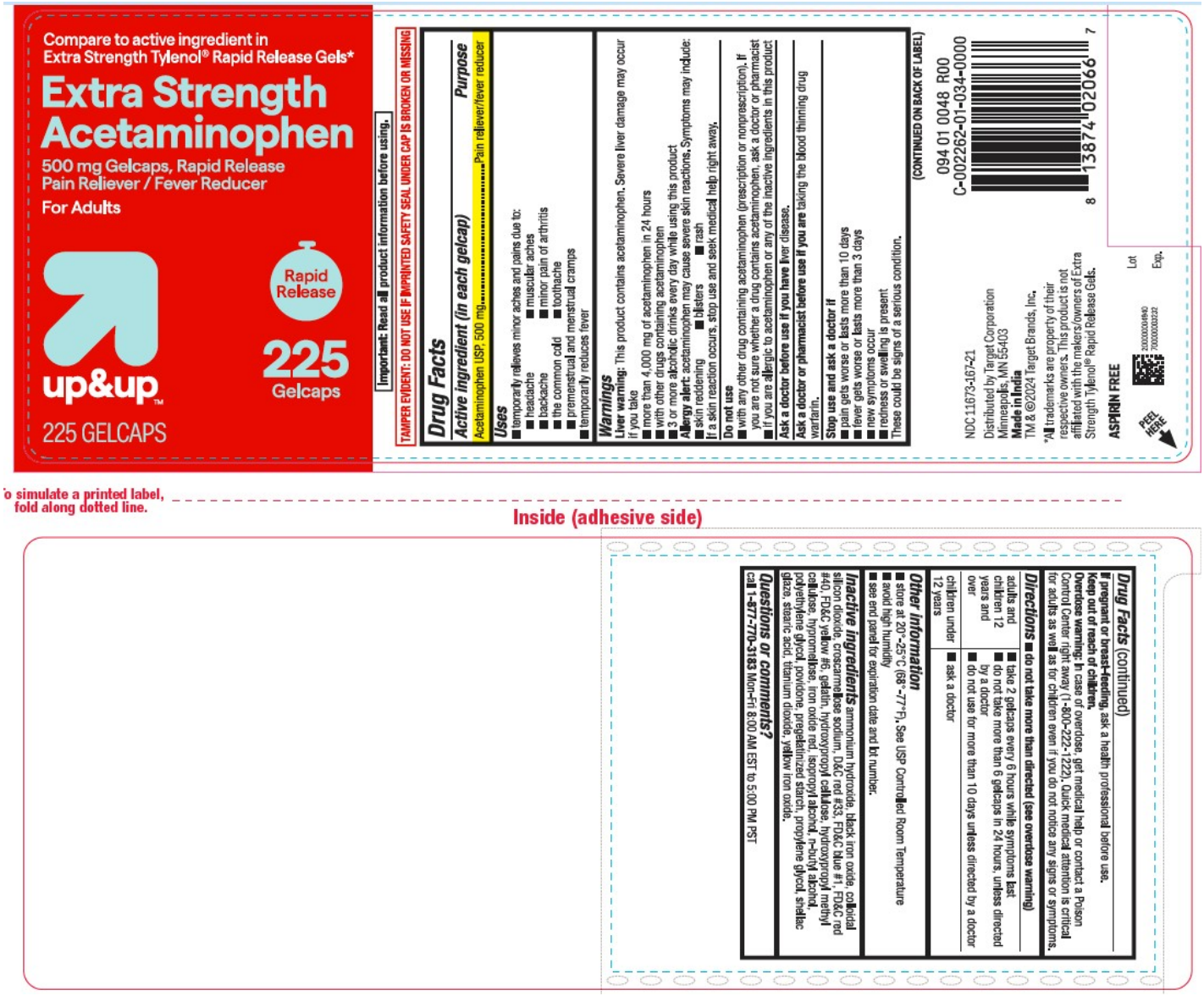

ACETAMINOPHEN

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-167
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**
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
AMMONIA (UNII: 5138Q19F1X)	
SHELLAC (UNII: 46N107B71O)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN (UNII: 2G86QN327L)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
BUTYL ALCOHOL (UNII: 8PJ61P6TS3)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
HYPROMELLOSE 2910 (3 MPA.S) (UNII: 0VUT3PMY82)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POVIDONE K30 (UNII: U725QWY32X)	
STARCH, CORN (UNII: O8232NY3SJ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
HYDROXYMETHYL CELLULOSE (UNII: 273FM27VK1)	

Product Characteristics

Color	gray (Encapsulated with red opaque and blue gray opaque hard gelatin shells)	**Score**	2 pieces
Shape	OVAL	**Size**	19mm
Flavor		**Imprint Code**	G1
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-167-05	50 in 1 BOTTLE; Type 0: Not a Combination Product	02/27/2023	
2	NDC:11673-167-10	100 in 1 BOTTLE; Type 0: Not a Combination Product	02/27/2023	
3	NDC:11673-167-21	225 in 1 BOTTLE; Type 0: Not a Combination Product	02/27/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	02/27/2023	

Labeler - TARGET CORPORATION (006961700)