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

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Drug Facts (continued)

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Drug Facts (continued)

Drug Facts (com next)

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Satisfaction guaranteed -Love it or your money back.

NDC 11673-167-05

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Extra Strength Acetaminophen

50 GELCAPS

മു 500 mg Gelcaps, Rapid Release Pain Reliever / Fever Reducer <u>rengtl</u>



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

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)

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Rapid Release Gels

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COATING FREE AREA



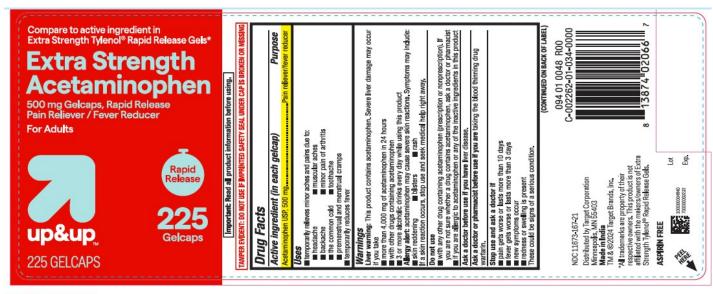
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COATING **FREE AREA**



Inside (adhesive side)

u**estions or comments?** 1-877-770-3183 Mon-Fri 8:00 AM EST to 5:00 PM PST. #1, FDSC red #40, FDSC yellow #5, gelatin, hydroxypropyl collubose, hydroxypropyl methyl cellubose, hyprometose, iron oxide red, isopropyl abohol, n-butyl alcohol, polyethylene glycol, povisione, pregelatinized starch, propylene glycol, shellec glaze, stearic acid, titanium dioxide, yellow iron ■ fever gets worse or lasts more than 3 days colloidal silicon dioxide, croscarmellose sodium, D&C red #33, FD&C blue 12 years and over adults and children do not take more than directed (see overdose warning) or symptoms. is critical for adults as well as for children even if you do not notice any signs Poison Control Center right away (1-800-222-1222). Quick medical attention new symptoms occu pain gets worse or lasts more than 10 days Drug Facts (continued) children under 12 years Stop use and ask a doctor if hinning drug warfarin. store at 20° - 25°C (68° - 77°F). See USP Controlled Room Temperature Other information nactive ingredients ammonium hydroxide, black iron oxide Mrections lavoid high humidity ep out of reach of children. pregnant or breast-feeding, ask a health professional before use. nese could be signs of a serious condition. redness or swelling is present see end panel for expiration date and lot number **erdose warning: h** case of overdose, get medical help or contact a k a doctor or pharmacist before use if you are taking the blood a doctor before use if you have liver disease do not take more than 6 gelcaps in 24 hours, take 2 gelcaps every 6 hours while symptoms ask a doctor do not use for more than 10 days unless unless directed by a doctor directed by a doctor



o simulate a printed label, fold along dotted line.

Inside (adhesive side)

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 ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN (UNIII - UNIII - UNIII

Inactive Ingredients	
Ingredient Name	Strength
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

FERRIC OXIDE YELLOW (UNII: EX43802MRT)	
AMMONIA (UNII: 5138Q19F1X)	
SHELLAC (UNII: 46N107B710)	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)	
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: 3WJQ0SDW1A)	
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			

P	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	

Revised: 11/2024 TARGET CORPORATION