ACETAMINOPHEN- acetaminophen tablet FAMILY DOLLAR

Extra Strength Pain Relief

Acetaminophen USP, 500mg Pain Reliever/Fever Reducer

FOR ADULTS

Active ingredient (in each caplet)

Acetaminophen USP, 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

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 use for more than 10 days unless directed by a doctor

children under

12 years **■** ask a doctor

Other information

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

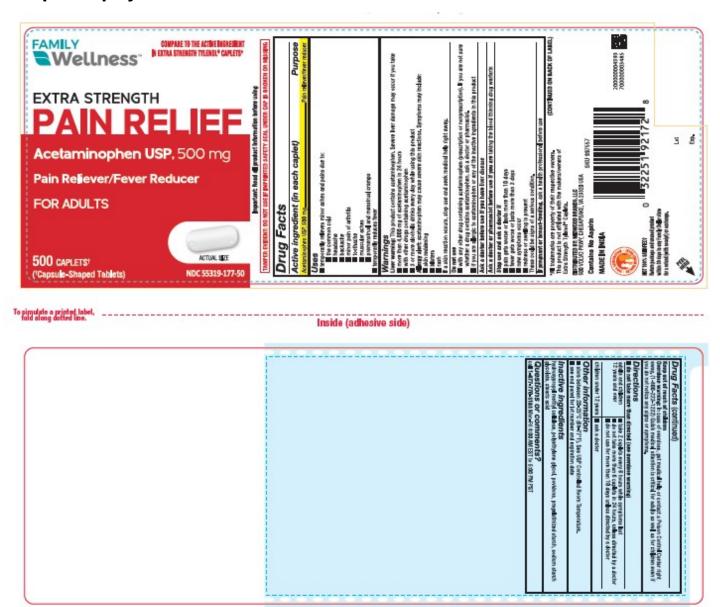
Inactive ingredients

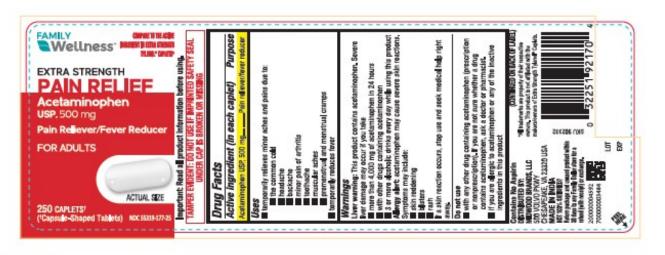
hydroxypropyl methyl cellulose, polyethylene glycol, povidone, pregelatinized starch, sodium starch glycolate, stearic acid

Questions or comments?

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

Pricipal Display Panel





To simulate a printed label, fold along dotted line.

Inside (adhesive side)

Drug Facts (continued)

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Children under

2 years

Other information

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ACETAMINOPHEN

acetaminophen tablet

Product Information				
	Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55319-177
	Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	500 mg		

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE K30 (UNII: U725QWY32X)	
STARCH, CORN (UNII: O8232NY3SJ)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics			
Color	white (WHITE TO OFF-WHITE)	Score	no score
Shape	CAPSULE	Size	17mm
Flavor		Imprint Code	G;551
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55319-177- 24	24 in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2023	
2	NDC:55319-177- 05	50 in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2023	
3	10	100 in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2023	
4	NDC:55319-177- 25	250 in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2023	
5	NDC:55319-177- 50	500 in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	09/01/2023	

Labeler - FAMILY DOLLAR (024472631)

Revised: 12/2023 FAMILY DOLLAR