

ACETAMINOPHEN- acetaminophen tablet, film coated
Sam's West Inc

Pain Reliever

Active ingredient (in each caplet)

Acetaminophen 500 mg

Purpose

Pain reliever/fever reducer

Uses

- temporarily relieves minor aches and pains due to:
 - headache
 - muscular aches
 - the common cold
 - backache
 - toothache
 - minor pain from arthritis
 - 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:

- rash
- skin reddening
- blisters

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.

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

- redness or swelling is present
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- any new symptoms appear

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away. Prompt 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**
- 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

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

Inactive ingredients

castor oil, hypromellose, povidone, sodium starch glycolate, starch, stearic acid

Questions or comments?

1-800-426-9391

Principal Display Panel

COMPARE TO **EXTRA STRENGTH**
TYLENOL® CAPLETS ACTIVE INGREDIENT*

M

**Member's
Mark™**

NDC 68196-031-06

EXTRA STRENGTH

Acetaminophen
500 mg CAPLETS

Pain Reliever/Fever Reducer

600 CAPLETS**

(**CAPSULE SHAPED TABLETS)

ACTUAL SIZE
FOR ADULTS

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

***This product is not manufactured or distributed by Kenvue Inc., owner of
the registered
trademark Extra Strength Tylenol® Caplets. 50844 ORG102417578**

DISTRIBUTED BY: SAM'S WEST, INC., BENTONVILLE, AR 72716



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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.
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 ■ redness or swelling is present
■ fever gets worse or lasts more than 3 days ■ any new symptoms appear
If pregnant or breast-feeding, ask a health professional before use.
Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Drug Facts (continued)

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Discard Seal,
Empty &
Replace Cap

**PLASTIC
BOTTLE**

how2recycle.info

members mark 44-175

ACETAMINOPHEN

acetaminophen tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68196-031
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		
CASTOR OIL (UNII: D5340Y2I9G)				
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)				
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)				
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)				
STARCH, CORN (UNII: O8232NY3SJ)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
Product Characteristics				
Color	white	Score	no score	
Shape	OVAL	Size	17mm	
Flavor		Imprint Code	44;175	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68196-031-06	600 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/14/2025	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013		11/14/2025	

Labeler - Sam's West Inc (051957769)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(68196-031)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(68196-031)

Establishment

Name	Address	ID/FEI	Business Operations
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LNK International, Inc.		832867894	manufacture(68196-031)
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Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(68196-031)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(68196-031)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117597853	pack(68196-031)