

**ACETAMINOPHEN EXTRA STRENGTH- acetaminophen tablet, coated**  
**Sam's West Inc**

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**Members Mark 44-519**

***Active ingredient (in each gelcap)***

Acetaminophen 500 mg

***Purpose***

Pain reliever/fever reducer

***Uses***

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

**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
- redness or swelling is present
- 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 gels every 6 hours while symptoms last
  - do not take more than 6 gels 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)
- avoid high humidity
- use by expiration date on package

***Inactive ingredients***

croscarmellose sodium, D&C red #33, FD&C blue #1, FD&C red #40, gelatin, hydroxypropyl cellulose, hypromellose, iron oxide black, iron oxide red, iron oxide yellow, polyethylene glycol, povidone, pregelatinized starch, propylene glycol, shellac glaze, stearic acid, titanium dioxide

***Principal Display Panel***

COMPARE TO **EXTRA  
STRENGTH TYLENOL®  
RAPID RELEASE GELS  
ACTIVE INGREDIENT\***

NDC 68196-919-05

**M**  
**Member's**  
**Mark™**

EXTRA STRENGTH  
**Acetaminophen**  
500 mg

Pain Reliever/Fever Reducer

- NON-ASPIRIN
- RELIEVES MINOR  
ACHES & PAINS

**400 GELCAPS**

ACTUAL SIZE

**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®  
Rapid Release Gels.  
50844 REV0525D51905

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

**PRODUCT OF CHINA**

**100% MONEY BACK GUARANTEE**  
**SUPERIOR QUALITY AND PERFORMANCE**

We would like to hear from you with any comments or suggestions.  
In the continental U.S. or Canada, you can call us at 1-800-426-9391  
from 8:30 a.m to 4:00 p.m. EST Monday-Friday.

<p><b>Drug Facts (continued)</b></p> <p>Stop use and ask a doctor if</p> <ul style="list-style-type: none"> <li>■ pain gets worse or lasts more than 10 days</li> <li>■ fever gets worse or lasts more than 3 days</li> <li>■ redness or swelling is present</li> <li>■ any new symptoms appear</li> </ul> <p>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.</p> <p><b>Directions</b> ■ do not take more than directed</p> <ul style="list-style-type: none"> <li>■ adults and children 12 years and over</li> <li>■ take 2 gelcaps every 6 hours while symptoms last</li> <li>■ do not take more than 6 gelcaps in 24 hours, unless directed by a doctor</li> <li>■ do not take for more than 10 days unless directed by a doctor</li> <li>■ children under 12 years: ask a doctor</li> </ul> <p><b>Other information</b> ■ store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)</p> <ul style="list-style-type: none"> <li>■ avoid high humidity</li> <li>■ use by expiration date on package</li> </ul> <p><b>Inactive ingredients</b> croscarmellose sodium, D&amp;C red #33, FD&amp;C blue #1, FD&amp;C red #40, gelatin, hydroxypropyl cellulose, hypromellose, iron oxide black, iron oxide red, iron oxide yellow, polyethylene glycol, povidone, pregelatinized starch, propylene glycol, shellac glaze, stearic acid, titanium dioxide</p>	<p>No Varnish/No print Lot no. &amp; Exp. Date</p>	<p>COMPARE TO EXTRA STRENGTH TYLENOL® RAPID RELEASE GELS ACTIVE INGREDIENT*</p> <p>NDC 68196-919-05</p> <p><b>Member's Mark™</b></p> <p>EXTRA STRENGTH <b>Acetaminophen</b> 500 mg</p> <p><b>Pain Reliever/Fever Reducer</b></p> <p><b>400 GELCAPS</b> ACTUAL SIZE</p> <ul style="list-style-type: none"> <li>• NON-ASPIRIN</li> <li>• RELIEVES MINOR ACHES &amp; PAINS</li> </ul>	<p>0 178742 109206 5</p> <p>*This product is not manufactured or distributed by Kenvue Inc., owner of the registered trademark Extra Strength Tylenol® Rapid Release Gels. 50844 REV0525D51905 DISTRIBUTED BY: SAM'S WEST, INC., BENTONVILLE, AR 72716 PRODUCT OF CHINA</p>	<p><b>Drug Facts</b> TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING</p> <table border="1"> <thead> <tr> <th>Active ingredient (in each gelcap)</th> <th>Purpose</th> </tr> </thead> <tbody> <tr> <td>Acetaminophen 500 mg</td> <td>Pain reliever/fever reducer</td> </tr> </tbody> </table> <p><b>Uses</b></p> <ul style="list-style-type: none"> <li>■ temporarily relieves minor aches and pains due to:             <ul style="list-style-type: none"> <li>■ headache</li> <li>■ the common cold</li> <li>■ backache</li> <li>■ minor pain from arthritis</li> <li>■ toothache</li> <li>■ muscular aches</li> <li>■ premenstrual and menstrual cramps</li> </ul> </li> <li>■ temporarily reduces fever</li> </ul> <p><b>Warnings</b> Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take</p> <ul style="list-style-type: none"> <li>■ more than 4,000 mg of acetaminophen in 24 hours</li> <li>■ with other drugs containing acetaminophen</li> <li>■ 3 or more alcoholic drinks every day while using this product</li> </ul> <p><b>Allergy alert:</b> Acetaminophen may cause severe skin reactions. Symptoms may include:</p> <ul style="list-style-type: none"> <li>■ skin reddening</li> <li>■ blisters</li> <li>■ rash</li> </ul> <p>If a skin reaction occurs, stop use and seek medical help right away.</p> <p>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.</p> <p>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.</p>	Active ingredient (in each gelcap)	Purpose	Acetaminophen 500 mg	Pain reliever/fever reducer
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Acetaminophen 500 mg	Pain reliever/fever reducer							

**Members Mark 44-519**

# ACETAMINOPHEN EXTRA STRENGTH

acetaminophen tablet, coated

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68196-919
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

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

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
D&C RED NO. 33 (UNII: 9DBA05BB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B71O)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

## Product Characteristics

<b>Color</b>	red, blue	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	19mm
<b>Flavor</b>		<b>Imprint Code</b>	L;5
<b>Contains</b>			

## Packaging

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:68196-919-05	400 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/10/2004	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	05/10/2004	

**Labeler** - Sam's West Inc (051957769)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	manufacture(68196-919) , pack(68196-919)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(68196-919)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(68196-919)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(68196-919)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(68196-919)

Revised: 8/2025

Sam's West Inc