

**TYLENOL EXTRA STRENGTH- acetaminophen capsule, liquid filled**  
**Kenvue Brands LLC**

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**TYLENOL® EXTRA STRENGTH**

***Drug Facts***

***Active ingredient (in each capsule)***

Acetaminophen 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**

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.**

### **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	<ul style="list-style-type: none"><li>▪ take 2 capsules every 6 hours while symptoms last</li><li>▪ do not take more than 6 capsules in 24 hours, unless directed by a doctor</li><li>▪ do not use for more than 10 days unless directed by a doctor</li></ul>
children under 12 years	ask a doctor

### **Other information**

- store between 20-25°C (68-77°F)
- **do not use if carton is opened. Do not use if foil inner seal imprinted with "TYLENOL" is broken or missing**

## **Inactive ingredients**

FD&C red no. 40, FD&C yellow no. 6, gelatin, glycerin, hypromellose, lecithin, medium chain triglycerides, polyethylene glycol, povidone, propylene glycol, sorbitol, titanium dioxide

## **Questions or comments?**

Call **1-877-895-3665** (toll-free) or **215-273-8755** (collect)

## **PRINCIPAL DISPLAY PANEL**

NDC 50580-469-18

***NEW***

***Extra Strength***

***TYLENOL***®

FOR ADULTS

Acetaminophen

Pain Reliever - Fever Reducer

***LIQUID GELS***

**Actual Size**

***18 Liquid Gels\****

***500 mg each***

**\*Liquid Filled Capsules**



22865003



**Important:** Read all product information before using. Keep this box for important information.

**Drug Facts**

**Active ingredient (in each capsule): Purpose**  
Acetaminophen 500 mg.....Pain reliever/fever reducer

**Uses**

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

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  - 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 redness
  - 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.

**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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- 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.

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**TYLENOL**<sup>®</sup>

**Acetaminophen**  
Pain Reliever-Fever Reducer

**LIQUID GELS**

\*Liquid Filled Capsules

Actual Size



**Ty 500**  
**18 Liquid Gels\***  
**500 mg each**

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Pat. www.kenvuepat.com  
www.tylenol.com

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**TYLENOL EXTRA STRENGTH**

acetaminophen capsule, liquid filled

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:50580-469
<b>Route of Administration</b>	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
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GLYCERIN (UNII: PDC6A3C0OX)	
GELATIN (UNII: 2G86QN327L)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SORBITOL (UNII: 506T60A25R)	

### Product Characteristics

Color	red	Score	no score
Shape	OVAL	Size	24mm
Flavor		Imprint Code	TY;500
Contains			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50580-469-18	1 in 1 CARTON	10/27/2025	
1		18 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:50580-469-40	1 in 1 CARTON	10/27/2025	
2		40 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:50580-469-75	1 in 1 CARTON	10/27/2025	
3		75 in 1 BOTTLE; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	10/27/2025	

