

**ACETAMINOPHEN- acetaminophen tablet**  
**Walmart**

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**Pain Reliever**  
**Acetaminophen USP, 500 mg**  
**Pain Reliever/Fever Reducer**  
**Contains No Aspirin**

**Active ingredient (in each gelcap)**

Acetaminophen USP, 500 mg

**Purpose**

Pain reliever/fever reducer

**Uses**

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

**Liver warning:**

This product contains acetaminophen. Severe liverdamage may occur if you take

- more than 4,000mg 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 seekmedical 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

- 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)
- avoid high humidity
- see bottom of the label 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, hypromellose, isopropyl alcohol, n-butyl alcohol, polyethylene glycol, povidone k-30, pregelatinized starch, propylene glycol, red iron oxide, shellac glaze, stearic acid, titanium dioxide, yellow iron oxide

## Questions or comments?

1-888-287-1915

**equate™** NDC 79903-269-40  
**EXTRA STRENGTH**  
**Pain Reliever**  
**Acetaminophen USP,**  
**500 mg**  
 Pain Reliever/Fever Reducer  
 Contains No Aspirin  
 Actual Size **500 mg EACH** **100 GELCAPS**

**Compare It To**  
 Take Strength  
 Take Well  
 Heat  
 Action  
 Action  
 Action

**IMPORTANT: READ THE DIRECTIONS AND WARNINGS BEFORE USE**  
**TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING**

**Drug Facts**  
**Active ingredient (in each gelcap)** Purpose  
 Acetaminophen USP, 500 mg 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

**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 redness ■ hives ■ rash  
 ■ 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.

**(CONTINUED ON BACK OF LABEL)**  
 634603  
 1-94346-26359-2  
 200000005373 7000000005956  
 LOT  
 EXP  
 Satisfaction guaranteed™ For questions or concerns call 1-888-287-1915  
 634603  
 Manufactured by:  
 Watson Pharmaceuticals, Inc.  
 10000 W. 16th Ave.  
 Golden, CO 80231  
**PRODUCT OF INDIA**  
 This product contains acetaminophen, the active ingredient in Tylenol. Do not use if the seal is broken or missing.  
 ♻️ **PLASTIC BOTTLE**  
 ♻️ **DO NOT REUSE**

## Inside (adhesive side)

**Drug Facts (continued)**  
 ■ If you are allergic to acetaminophen or any of the inactive ingredients in this product.  
 Ask a doctor or pharmacist before use if you have liver disease, kidney or lung problems.  
 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, 1-800-272-1222. 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 with symptoms that 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) ■ avoid high humidity  
 ■ see bottom of the bottle 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, hypromellose, isopropyl alcohol, n-butyl alcohol, polyethylene glycol, povidone k-30, pregelatinized starch, propylene glycol, red iron oxide, shellac glaze, stearic acid, titanium dioxide, yellow iron oxide

**Questions or comments?**  
 1-888-287-1915

**equate™**  
**EXTRA STRENGTH**  
**Pain Reliever**  
**Acetaminophen USP,**  
**500 mg**  
 Compare to Extra Strength Tylenol® Rapid Release Gels active ingredient\*  
 NDC 79903-269-22  
 Actual Size  
 Pain Reliever/Fever Reducer  
 Contains No Aspirin  
 500 mg EACH  
 225 GELCAPS  
**NOT FOR HOUSEHOLDS WITH YOUNG CHILDREN**  
**IMPORTANT: READ THE DIRECTIONS AND WARNINGS BEFORE USE**  
**TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING**

**Drug Facts**  
**Active ingredient (in each gelcap)**  
 Acetaminophen USP, 500 mg— 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  
 ■ menstrual 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 redness ■ hives  
 ■ 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

Satisfaction guaranteed—For questions or comments please call 1-888-387-4815.  
 \*As compared to Extra Strength Tylenol® Rapid Release Gels.  
 LOT: 2000000065372  
 PRODUCT OF INDIA  
 This product is not manufactured or distributed by the name of the manufacturer listed below.  
 Manufactured by:  
 ■ How2Recycle.com  
 ■ 2000000065372  
 ■ 7000000039925  
 ■ 19434626360  
 ■ 8  
 Lot Exp.  
 PLASTIC BOTTLE  
 Recycle with plastic bottles

(CONTINUED ON BACK OF LABEL)

Inside (adhesive side)

**Drug Facts (continued)**  
 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: 1-800-272-2273. 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 ■ take 2 gelcaps every 6 hours while symptoms last  
 ■ do not take more than 2 gelcaps in 24 hours, unless directed by a doctor  
 7 years and over ■ 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)  
 ■ avoid high humidity  
 ■ see bottom of the label for expiration date and lot number

**Inactive ingredients**  
 acetaminophen hydrochloride, thick-film coating: calcium stearate, croscarmellose sodium, D1-C, D1-E3, FDAC, K1, FDAC, K1 and K2, FDAC, K1 and K2, gelatin, hydroxypropyl methylcellulose, hydroxypropyl starch, polyvinyl alcohol, polyethylene glycol, polyethylene glycol 400, croscarmellose sodium, polyvinyl alcohol, red iron oxide, silicon dioxide, stearic acid, titanium dioxide, yellow iron oxide

**Questions or comments?**  
 1-888-387-4815

**ACETAMINOPHEN**

acetaminophen tablet

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:79903-269
<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
AMMONIA (UNII: 5138Q19F1X)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	

<b>D&amp;C RED NO. 33</b> (UNII: 9DBA05BB0L)
<b>FERROSFERRIC OXIDE</b> (UNII: XM0M87F357)
<b>GELATIN</b> (UNII: 2G86QN327L)
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)
<b>BUTYL ALCOHOL</b> (UNII: 8PJ61P6TS3)
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)
<b>POVIDONE K30</b> (UNII: U725QWY32X)
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)
<b>ISOPROPYL ALCOHOL</b> (UNII: ND2M416302)
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)
<b>FERRIC OXIDE YELLOW</b> (UNII: EX438O2MRT)
<b>SHELLAC</b> (UNII: 46N107B71O)
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)

### Product Characteristics

<b>Color</b>	red (red, blue (blue-gray))	<b>Score</b>	no score
<b>Shape</b>	CAPSULE (banded two-part capsule)	<b>Size</b>	19mm
<b>Flavor</b>		<b>Imprint Code</b>	G;1
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-269-10	100 in 1 BOTTLE; Type 0: Not a Combination Product	01/16/2025	
2	NDC:79903-269-22	225 in 1 BOTTLE; Type 0: Not a Combination Product	01/16/2025	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	01/16/2025	

**Labeler** - Walmart (051957769)