

**ACETAMINOPHEN- acetaminophen tablet**  
**AARNA USA INC.**

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**ACTIVE INGREDIENT (IN EACH TABLET)**

Acetaminophen 500 mg

**PURPOSE**

Pain Reliever/Fever Reducer

**USES**

To reduce fever and for the temporary relief of 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
- 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 non-prescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or

pharmacist.

- if the user has ever had an allergic reaction to this product or any of its ingredients.
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**Ask a doctor before use if the user has**

has liver disease

**Ask a doctor or pharmacist before use if the user is 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:**

Taking more than the recommended dose (overdose) may cause liver damage. 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)**

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	◦ take 2 tablets every 6 hours while symptoms last
	◦ do not take more than 6 tablets in 24 hours, unless directed by a doctor
adults & children 12 years and over	◦ do not use for more than 10 days unless directed by a doctor
children under 12 years	◦ ask a doctor

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**OTHER INFORMATION**

store at 20° to 25°C (68° to 77°F)

**INACTIVE INGREDIENTS:**

Gelatinized starch, magnesium stearate, povidone

Leland, NC 28451

\* This product is not manufactured or distributed by Johnson and Johnson, consumer inc., distributor of regular Tylenol Tablets.

[illegible]

N 82566 000218

NDC 82568-0002-2

# Acetaminophen 500MG TABLETS

**EXTRA STRENGTH**

**ASPIRIN FREE  
STRONG PAIN &  
FEVER RELIEVER**

\*Compare to other Ingredient  
in Regular Strength Tylenol® Tablets

DO NOT USE WITH  
OTHER MEDICINES  
CONTAINING  
ACETAMINOPHEN

**1000  
TABLETS**

## Drug Facts

### Active ingredient (in each tablet)

**Acetaminophen 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
  - 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 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 have ever had an allergic reaction to this product or any of the ingredients.

### Ask a doctor before use if the user

- has liver disease

### Ask a doctor or pharmacist before use if the user is

taking the blood thinning drug warfarin

Child-Resistant Packaging

## Drug Facts (continued)

### 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
- any 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:** Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center (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 & children 16 years and over

- take 2 tablets every 6 hours while symptoms last
- do not take more than 6 tablets in 24 hours, unless directed by a doctor

do not use for more than 10 days unless directed by a doctor

#### child under 12 years

ask a doctor

**Other information:** store at 20° to 25°C (68° to 77°F)

**Inactive ingredients:** gelatinized starch, magnesium stearate, povidone.

**Questions or comments?** Call toll-free 1-877-225-6999

Do not use if imprinted safety seal under cap is broken or missing

\*This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Tylenol® Extra Strength

**Adamo USA**  
Manufactured for:  
**ADAMO USA, Inc.**  
Lancaster, NC 28645

**No Varnish**

## acetaminophen tablet

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82568-0002
Route of Administration	ORAL		

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

Ingredient Name	Strength
MAGNESIUM STEARATE (UNII: 70097M6I30)	
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE (UNII: FZ989GH94E)	

<b>Color</b>	white	<b>Score</b>	2 pieces
<b>Shape</b>	ROUND	<b>Size</b>	5mm
<b>Flavor</b>		<b>Imprint Code</b>	A500
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82568-0002-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2025	
2	NDC:82568-0002-2	1000 in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2025	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	05/15/2025	

**Labeler** - AARNA USA INC. (118515992)

Revised: 5/2025

AARNA USA INC.