

ACETAMINOPHEN- acetaminophen tablet
Akron Pharma Inc.

Acetaminophen 500 mg Tablets
Extra Strength

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

- adult takes more than 4,000 mg of acetaminophen in 24 hours
- child takes more than 5 doses in 24 hours
- taken with other drugs containing acetaminophen
- adult has 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.

Ask a doctor before use if the user has

- has liver disease
- is a child with pain of arthritis

Ask a doctor or pharmacist before use if your child is taking the blood thinning drug warfarin

Stop use and ask a doctor if:

- pain gets worse or lasts more than 10 days (for adults) or 5 days (for children)
- 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)

adults & children 12 years and over	<ul style="list-style-type: none">• 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
children under 12 years	<ul style="list-style-type: none">• 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

Manufactured for
Akron Pharma, Inc.,

373 RT US46 W Building E,

Suite 117, Fairfield, NJ - 07004

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

Drug Facts
Active ingredient (in each tablet) Acetaminophen 500 mg Pain reliever/ Fever reducer
Uses ■ temporarily relieves minor aches and pains due to: ■ headache ■ muscle 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 the user has ever had an allergic reaction to this product or any of its 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 under label) 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 N07134005-01 <div style="border: 1px solid black; padding: 5px; text-align: center; color: red; font-weight: bold;">No Tamish</div> <div style="font-size: 8px; text-align: right;"> PEEL HERE FOR MORE DRUG FACTS </div>

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 ■ new symptoms occur ■ redness or swelling is present.
 These could be signs of a serious condition.

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Keep out of reach of children.

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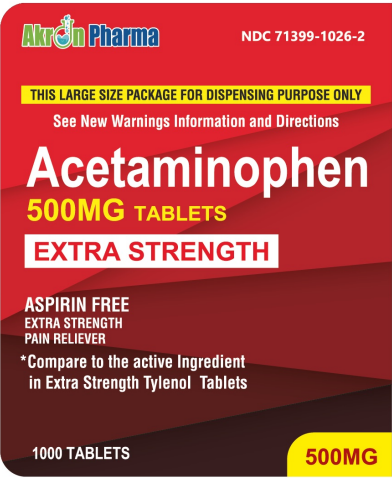
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Children under 12 years:
 ask a doctor

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Acetaminophen
500MG TABLETS
EXTRA STRENGTH

1000 TABLETS

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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)

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No Varnish

ACETAMINOPHEN acetaminophen tablet				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71399-1026	
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	
STARCH, CORN (UNII: O8232NY3SJ)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
POVIDONE (UNII: FZ989GH94E)				
Product Characteristics				
Color	white	Score	2 pieces	
Shape	ROUND	Size	5mm	
Flavor		Imprint Code	A500	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-1026-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	12/06/2024	
2	NDC:71399-1026-2	1000 in 1 BOTTLE; Type 0: Not a Combination Product	12/06/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	12/06/2024	

Labeler - Akron Pharma Inc. (067878881)

Revised: 12/2024

Akron Pharma Inc.