ACETAMINOPHEN- acetaminophen tablet Major Pharmaceuticals

5427-Major

Drug Facts

Active Ingredient (in each tablet)

Acetaminophen 500 mg

Purpose

Pain reliever/ Fever reducer

Uses

Temporarily reduces fever and relieves minor aches and pains due to:

- headache
- muscular aches
- backache
- minor pain of arthritis
- the common cold
- toothache
- premenstrual and menstrual cramps

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.

- 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 \square

- 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 tablets every 6 hours while symptoms last
- do not take more than 6 tablets 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 in a dry place at 15° - 30°C (59° - 86°F).

Povidone, pregelatinized starch, sodium starch glycolate, stearic acid

Questions or comments?

Call 1-800-231-4670

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Distributed by:

MAJOR® PHARMACEUTICALS Indianapolis, IN 46268

*This product is not manufactured or distributed by Kenvue Inc., owner of the registered trademark Tylenol®.

MAJOR

THIS LARGE SIZE PACKAGE FOR DISPENSING PURPOSE ONLY

See New Warnings Information and Directions

Acetaminophen Extra Strength Tablets

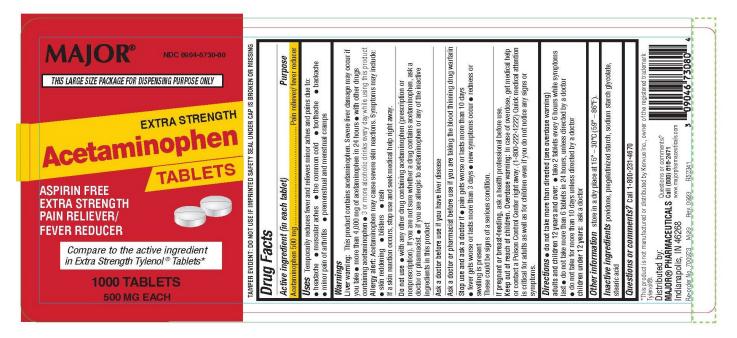
Aspirin Free

Extra Strenght Pain reliever

Compare to active ingredient in Extra Strength Tylenol®* Tablets

1000 tablets

500mg each



ACETAMINOPHEN

acetaminophen tablet

Product	Information	

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0904-6730

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)

ACETAMINOPHEN (UNII: 36209ITL9D) ACETAMINOPHEN 500 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
POVIDONE K30 (UNII: U725QWY32X)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Floduct Characteristics					
Color	white	Score	2 pieces		
Shape	ROUND	Size	12mm		
Flavor		Imprint Code	54;27		

Contains

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904- 6730-60	100 in 1 BOTTLE; Type 0: Not a Combination Product	09/12/2018	
2	NDC:0904- 6730-80	1000 in 1 BOTTLE; Type 0: Not a Combination Product	09/12/2018	
3	NDC:0904- 6730-59	1 in 1 CARTON	09/12/2018	
3		100 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:0904- 6730-61	10 in 1 BOX, UNIT-DOSE	09/12/2018	
4		10 in 1 BLISTER PACK; 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	09/12/2018		

Labeler - Major Pharmaceuticals (191427277)

Revised: 5/2024 Major Pharmaceuticals