ACETAMINOPHEN- acetaminophen tablet
Cardinal Health

5427-Cardinal

Drug Facts

Active ingredient

Acetaminophen 500 mg

Purpose

Pain Reliever/ Fever Reducer

Uses

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

● headache

● muscular aches

● common cold

● toothache

● backache

● minor pain of arthritis

● 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.

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 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 at room temperature

Povidone, pregelatinized starch, sodium starch glycolate, stearic acid

Questions or comments?

Call (800) 231-4670

*This product is not manufactured or distributed by McNeil Consumer Healthcare, owner of the registered trademark Tylenol® Extra Strength.

DISTRIBUTED BY CARDINAL HEALTH, DUBLIN, OHIO 43017

www.myleader.com 1-800-200-6313

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

100% Money Back Guarantee

Return to place of purchase if not satisfied.

LEADER TM

NDC 70000-0410-2

Extra Strength **Acetaminophen**

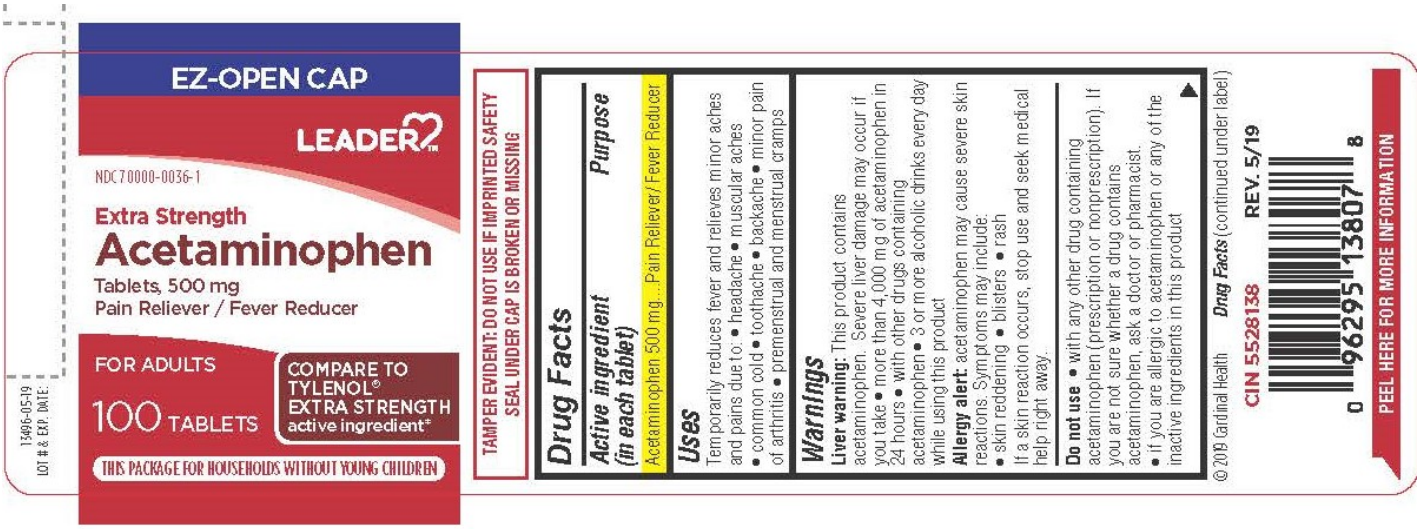
Tablets, 500 mg

Pain Reliever/ Fever Reducer

For Adults

Compare to Tylenol ® Extra Strength active ingredient*

100 Tablets



ACETAMINOPHEN

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70000-0410
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**
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

STEARIC ACID (UNII: 4ELV7Z65AP)	
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE K30 (UNII: U725QWY32X)	

Product 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:70000-0410-2	100 in 1 BOTTLE; Type 0: Not a Combination Product	11/05/2018	
2	NDC:70000-0410-1	1 in 1 CARTON	12/19/2018	
2		60 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	11/05/2018	

ACETAMINOPHEN

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70000-0036
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
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE K30 (UNII: U725QWY32X)	

Product 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:70000-0036-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/28/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	05/28/2019	

Labeler - Cardinal Health (063997360)

Revised: 1/2024 Cardinal Health