ACETAMINOPHEN- acetaminophen tablet A-S Medication Solutions

Major Pharmaceuticals Acetaminophen Drug Facts

Active ingredient (in each caplet)

Acetaminophen 500 mg

Purpose

Pain reliever/fever reducer

Uses

- temporarily relieves minor aches and pains due to:
- the common cold
- headache
- backache
- · minor pain of arthritis
- toothache
- muscular aches
- 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 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 its ingredients

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 caplets every 6 hours while symptoms last do not take more than 6 caplets 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			

Inactive ingredients

carnauba wax, corn starch*, croscarmellose sodium*, hypromellose, polyethylene glycol, povidone, pregelatinized starch, sodium starch glycolate*, stearic acid

*may contain one or more of these ingredients

Questions or comments?

1-800-616-2471

HOW SUPPLIED

Product: 50090-5272

NDC: 50090-5272-0 40 TABLET in a BOTTLE

NDC: 50090-5272-1 100 TABLET in a BOTTLE

ACETAMINOPHEN TABLET



ACETAMINOPHEN

acetaminophen tablet

Product Information	
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:50090-5272(NDC:0904-6720)

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN 500 mg

Inactive	Ingredients

Ingredient Name	Strength		
CARNAUBA WAX (UNII: R12CBM0EIZ)			
STARCH, CORN (UNII: O8232NY3SJ)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			

POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)

STEARIC ACID (UNII: 4ELV7Z 65AP)

CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)

Product Characteristics			
Color	WHITE	Score	no score
Shape	OVAL	Size	16mm
Flavor		Imprint Code	L484
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:50090- 5272-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	02/26/2021		
2	NDC:50090- 5272-0	40 in 1 BOTTLE; Type 0: Not a Combination Product	10/19/2020		

Marketing Information			
Marketing Category	• • • • • • • • • • • • • • • • • • • •		Marketing End Date
OTC Monograph Drug	M013	07/26/2018	

Labeler - A-S Medication Solutions (830016429)

Establishment				
Name	Address	ID/FEI	Business Operations	
A-S Medication Solutions		830016429	RELABEL(50090-5272), REPACK(50090-5272)	

Revised: 3/2024 A-S Medication Solutions