ACETAMINOPHEN- acetaminophen tablet Kroger Company

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Kroger Co. 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					

Other information

• store at 20-25°C (68-77°F)

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

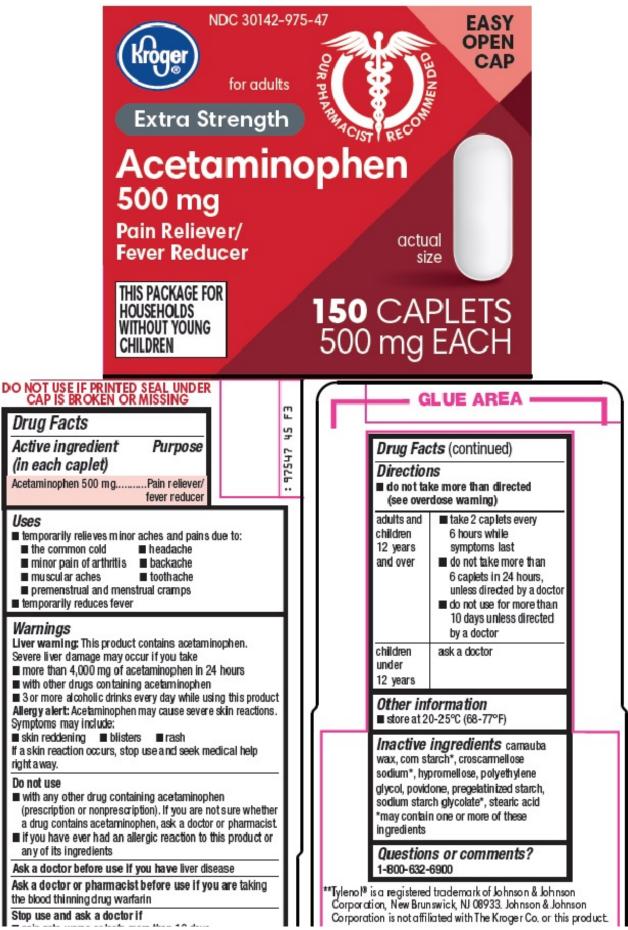
Questions or comments?

1-800-632-6900

Principal Display Panel

COMPARE TO the active ingredient of EXTRA STRENGTH TYLENOL® CAPLETS See back panel THIS PACKAGE FOR HOUSEHOLDS WITHOUT YOUNG CHILDREN EASY OPEN CAP OUR PHARAMCIST RECOMMENDED for adults Extra Strength Acetaminophen 500 mg Pain Reliever/Fever Reducer actual size THIS PACKAGE FOR HOUSEHOLDS WITHOUT YOUNG CHILDREN 150 CAPLETS 500 mg EACH

COMPARE TO the active ingredient of EXTRA STRENGTH TYLENOL® CAPLETS **See back panel





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acetaminophen tablet							
Product Information							
Product Type	HUMAN OT	C DRUG	Item Code (Source) ND		NDC:301	DC:30142-975	
Route of Administration	ORAL						
Active Ingredient/Activ	e Moiety						
In	gredient Na	ame		Basis of S	Strength	Strengt	
	CETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII: 36209ITL9D) ACETAMINOP				-	500 mg	
Inactive Ingredients	Ingrod	liant Nama			c	trongth	
Ingredient Name						trength	
•							
STARCH, CORN (UNII: 08232N)	′3SJ)						
STARCH, CORN (UNII: 08232N) HYPROMELLOSE, UNSPECIFIE	(UNII: 3NXW		A)				
STARCH, CORN (UNII: 08232N) HYPROMELLOSE, UNSPECIFIE POLYETHYLENE GLYCOL, UNS	'3SJ) Ed (UNII: 3NXW Specified (UN	NII: 3WJQ0SDW1	А)				
STARCH, CORN (UNII: 08232N) HYPROMELLOSE, UNSPECIFIE POLYETHYLENE GLYCOL, UNS POVIDONE, UNSPECIFIED (UN	73SJ) ED (UNII: 3NXW SPECIFIED (UN III: FZ989GH941	NII: 3WJQ0SDW1	A)				
CARNAUBA WAX (UNII: R12CBM STARCH, CORN (UNII: 08232N) HYPROMELLOSE, UNSPECIFIE POLYETHYLENE GLYCOL, UN POVIDONE, UNSPECIFIED (UN STEARIC ACID (UNII: 4ELV7Z65 CROSCARMELLOSE SODIUM (73SJ) ED (UNII: 3NXW SPECIFIED (UN III: FZ989GH941 AP)	NII: 3MJQOSDW1. E)	Α)				
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STARCH, CORN (UNII: 08232N) HYPROMELLOSE, UNSPECIFIE POLYETHYLENE GLYCOL, UNS POVIDONE, UNSPECIFIED (UN STEARIC ACID (UNII: 4ELV7Z65 CROSCARMELLOSE SODIUM (Product Characteristic Color	(3SJ) ED (UNII: 3NXW) SPECIFIED (UN III: FZ989GH941 AP) UNII: M28OL1H SS WHITE	NII: 3WJQ0SDW1 E) H48) Score Size		•	16mm		
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#	ltem Code	Package Description	Marketing Start Date	Marketing End Date					
1	NDC:30142-975- 47	150 in 1 BOTTLE; Type 0: Not a Combination Product	04/05/2014						
M	Marketing Information								
	Marketing	Application Number or Monograph	Marketing Start	Marketing End					
	Category	Citation	Date	Date					
OT fin	ر کی C monograph not		Date 04/05/2014	Date					

Labeler - Kroger Company (006999528)

Revised: 11/2022

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