TOPCARE PAIN RELIEF- acetaminophen tablet Praxis, LLC

Topcare Health Pain Relief 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
 - muscular aches
 - toothache
 - o 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

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
- new symptoms occur
- fever gets worse or lasts more than 3 days
- 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 vears 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?

1-888-423-0139

Principal Display Panel

TopCare ®health

COMPARE TO EXTRA STRENGTH TYLENOL ® CAPLETS ACTIVE INGREDIENT

EXTRA STRENGTH

Pain Relief

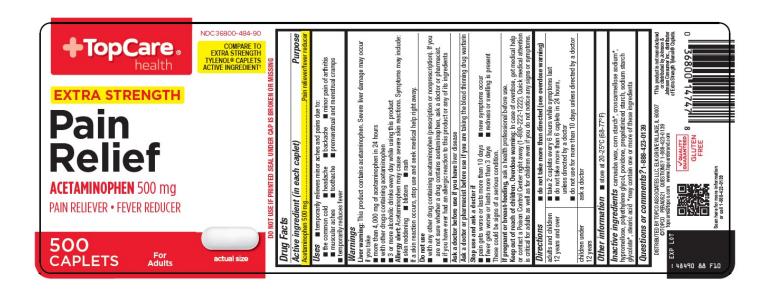
ACETAMINOPHEN 500 mg

PAIN RELIEVER • FEVER REDUCER

500 CAPLETS

For Adults

actual size



TOPCARE PAIN RELIEF

acetaminophen tablet

Product	Information
Product	iniormation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:59368-210

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN 500 mg

Inactive Ingredients

Ingredient Name	Strength
CARNAUBA WAX (UNII: R12CBM0EIZ)	
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

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

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:59368-210- 07	2 in 1 PACKAGE	07/15/1987		
1		500 in 1 BOTTLE; Type 0: Not a Combination Product			
2	NDC:59368-210- 06	500 in 1 BOTTLE; Type 0: Not a Combination Product	07/15/1987		
3	NDC:59368-210- 03	1 in 1 CARTON	07/15/1987		
3		100 in 1 BOTTLE; Type 0: Not a Combination Product			
4	NDC:59368-210- 04	1 in 1 CARTON	07/15/1987		
4		50 in 1 BOTTLE; Type 0: Not a Combination Product			
5	NDC:59368-210- 05	2 in 1 CARTON	07/15/1987		
5		100 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	07/15/1987	

Labeler - Praxis, LLC (016329513)

Establis	hment		
Name	Address	ID/FEI	Business Operations

Revised: 1/2023 Praxis, LLC