PAIN RELIEF EXTRA STRENGTH- acetaminophen tablet CVS Pharmacy

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.

CVS Pharmacy, Inc. 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
- toothache
- muscular aches
- premenstrual and menstrual cramps
- temporarily reduces fever

Warnings

Liver warning: This product contains acetaminophen. The maximum daily dose of this product is 6 caplets (3,000 mg) in 24 hours. 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

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.

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 Liver 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?

1-800-719-9260

Principal Display Panel

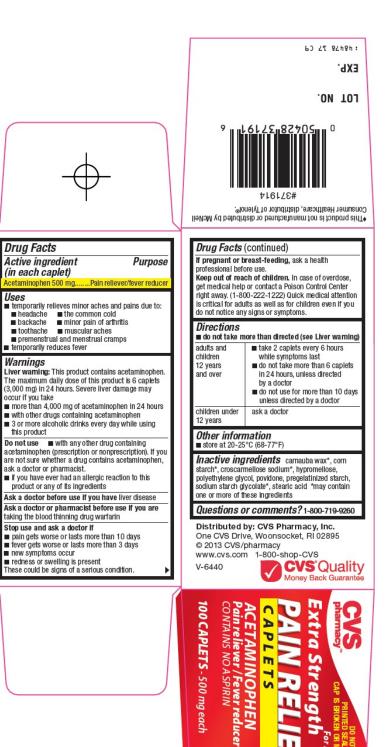
Compare to the active ingredient in Tylenol®

For Adults

Extra Strength

PAIN RELIEF CAPLETS
ACETAMINOPHEN
Pain reliever / Fever reducer
actual size
CONTAINS NO ASPIRIN
100 CAPLETS
500 mg each





PAIN RELIEF EXTRA STRENGTH

acetaminophen tablet

Drug Facts

Uses

Warnings

this product

Active ingredient

temporarily reduces fever

(in each caplet)

Product Information			
Product Type	HUMAN OTC DRUG LABEL	Item Code (Source)	NDC:59779-484
Route of Administration	ORAL	DEA Sche dule	

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINO PHEN (ACETAMINO PHEN)	ACETAMINOPHEN	500 mg

Inactive Ingredients				
Ingredient Name	Strength			
CARNAUBA WAX				
HYPROMELLOSES				
POLYETHYLENE GLYCOLS				
POVIDONES				
STEARIC ACID				
STARCH, CORN				
CROSCARMELLOSE SODIUM				

Product Characteristics				
Color	WHITE	Score	no score	
Shape	CAPSULE (caplet)	Size	16 mm	
Flavor		Imprint Code	L484	
Contains				

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1 1	NDC:59779-484-71	1 in 1 CARTON		
1		50 in 1 BOTTLE		
2 1	NDC:59779-484-78	1 in 1 CARTON		
2		100 in 1 BOTTLE		
3 1	NDC:59779-484-85	1 in 1 CARTON		
3		250 in 1 BOTTLE		
4 1	NDC:59779-484-62	1 in 1 CARTON		
4		24 in 1 BOTTLE		
5 1	NDC:59779-484-76	1 in 1 CARTON		
5		120 in 1 BOTTLE		
6 1	NDC:59779-484-72	14 in 1 CARTON		
6		60 in 1 BOTTLE		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	07/15/1991	

Labeler - CVS Pharmacy (062312574)

Revised: 11/2013 CVS Pharmacy