#### FAMILY WELLNESS PAIN RELIEF- acetaminophen tablet Family Dollar Services Inc

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.

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# Family Dollar Services, 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. 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	<ul> <li>take 2 caplets every 6 hours while symptoms last</li> <li>do not take more than 6 caplets in 24 hours, unless directed by a doctor</li> <li>do not use for more than 10 days unless directed by a doctor</li> </ul>
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

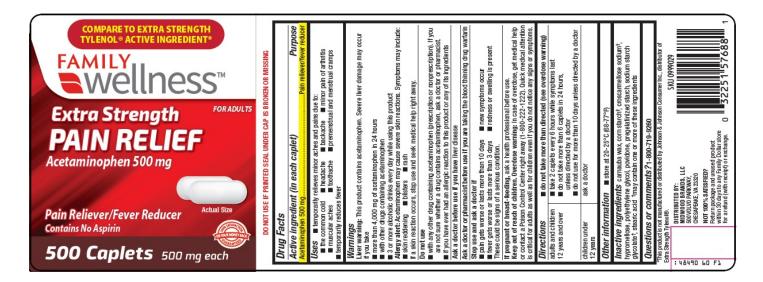
# Questions or comments?

1-800-719-9260

## **Principal Display Panel**

COMPARE TO EXTRA STRENGTH TYLENOL® ACTIVE INGREDIENT FOR ADULTS Extra Strength PAIN RELIEF Acetaminophen 500 mg Actual Size Pain Reliever/Fever Reducer Contains No Aspirin 100% SATISFACTION GUARANTEED OR YOUR MONEY BACK 500 Caplets

500 mg each



#### FAMILY WELLNESS PAIN RELIEF acetaminophen tablet **Product Information** HUMAN OTC DRUG **Product Type** Item Code (Source) NDC:55319-739 **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, UNSPI	HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)										
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)											
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)											
STEARIC ACID (UNII: 4ELV7Z65AP)											
CROSCARMELLOSE SO											
Product Characteristics											
Color	WHITE	Score	T	io score							
	OVAL										
Shape	UVAL	Size		6 mm							
Flavor		Imprint Code	1	L484							
Contains											
Packaging											
# Item Code	Package Desci	iption Marketing Start Dat		Marketing End Date							
<b>1</b> NDC:55319-739-90 500	0 in 1 BOTTLE; Type 0: Not a	Combination Product	02/26/2020								
Marketing Information											
Marketing Category	Application Number or	Monograph Citation	Marketing Start Date	e Marketing End Date							
OTC monograph not final	2 monograph not final part343		02/26/2020								

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Revised: 3/2020

Family Dollar Services Inc