# PAIN RELIEVER- acetaminophen 500 mg, film-coated tablet tablet, film coated

Lil' Drug Store Products, Inc.

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**Pain Reliever** 

# **Drug Facts**

# Active ingredient (in each caplet)

Acetaminophen USP 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**

**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

**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

#### 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 are allergic to acetaminophen or any of the inactive ingredients in this product

# 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

## 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**

**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**

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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

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store between 20° to 25°C (68° to 77°F)

## **Inactive ingredients**

Inactive ingredients povidone, pregelatinized starch (maize), stearic acid

## Questions or comments?

Call toll-free 1-877-507-6516 (M-F 8AM-4:30PM CST)

\*\*This product is not manufactured or distributed by JOHNSON & JOHNSON CONSUMER, INC., McNeil Consumer Healthcare Division, distributor of Tylenol® Extra Strength Caplets.

## PDP/Package

[caduceus]

Compare to the Active Ingredient in Tylenol® Extra Strength Caplets\*\*

Extra Strength Pain Reliever

Acetaminophen 500 mg Pain Reliever/Fever Reducer

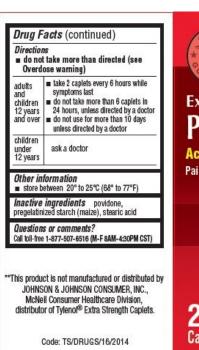
[caplet image]

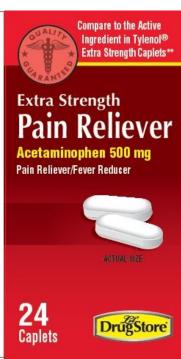
**ACTUAL SIZE** 

24

# **Caplets**

[Lil' Drug Store logo]





TAMPER EVIDENT: DO NOT USE IF CARTON IS OPEN OR IF Printed foil seal under cap is torn or missing.

WARNINGS AND PRODUCT INFORMATION

#### Drug Facts

Active ingredient (in each caplet)

Purpose

Acetaminophen USP 500 mg......Pain reliever.

- 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

#### Drug Facts (continued)

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 are allergic to acetaminophen or any of the inactive ingredients in this product

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 sings or symptoms. signs or symptoms.

## **PAIN RELIEVER**

acetaminophen 500 mg, film-coated tablet tablet, film coated

#### **Product Information**

**Product Type** HUMAN OTC DRUG **Item Code (Source)** NDC:66715-6867

**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**

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Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE K30 (UNII: U725QWY32X)	
PALMITOSTEARIC ACID (UNII: 08Y7S3B85M)	

#### **Product Characteristics**

Color	white (white to off-white)	Score	no score	
Shape	OVAL (capsule-shaped biconvex)	Size	18mm	
Flavor		Imprint Code	N79	
Contains				

#### **Packaging**

#	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66715- 6867-4	1 in 1 CARTON	03/04/2022	
1		24 in 1 BOTTLE, PLASTIC; 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	03/04/2022			

# **Labeler -** Lil' Drug Store Products, Inc. (093103646)

Revised: 12/2023 Lil' Drug Store Products, Inc.