LIL DRUG STORE PAIN RELIEVER EXTRA STRENGTH- acetaminophen tablet Lil' Drug Store Products, 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.

Lil' Drug Store ® Pain Reliever

Extra Strength

Drug Facts

Active ingredient (in each tablet)

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 tablets (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.

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 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 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

Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. 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 tablets every 6 hours while symptoms last do not take more than 6 tablets 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

- do not use if vial is open or damaged or safety seal has been removed
- store at room temperature 59-86°F (15-30°C)

Inactive ingredients

corn starch, hypromellose, maltodextrin 1 , microcrystalline cellulose 1 , polyethylene glycol, povidone 1 , pregelatinized starch 1 , sodium starch glycolate 1 , stearic acid, titanium dioxide 1

1 may contain

Questions or comments?

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

PRINCIPAL DISPLAY PANEL - 10 Tablet Vial Label

★ QUALITY GUARANTEED ★

Compare to the Active Ingredient in Tylenol® Extra Strength*

Extra Strength Pain Reliever

Acetaminophen, 500 mg Pain Reliever/Fever Reducer

10 Tablets

Lil'

DrugStore ®

OPEN HERE FOR DRUG FACTS INFORMATION

Lil' Drug Store Products, Inc. does not own the Tylenol® Extra Strength trademark. McNeil Consumer Healthcare Division of McNeil-PPC, Inc. *This product is not manufactured or distributed by

> Cedar Rapids, IA 52404 9300 Earhart Lane SW Product manufactured for: Lif Drug Store Products, Inc.

(M-F 8 AM-4:30PM CST) call toll-free 1-877-507-6516 Questions or comments?

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Drug Facts (continued)



Compare to the Active Ingredient in Tylenol® Extra Strength*

Extra Strength
Pain Reliever

Acetaminophen, 500 mg Pain Reliever/Fever Reducer

Tablets



Drug Facts Active ingredient (in each tablet)

Purpose

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Uses

SAFETY SEALED

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LIL DRUG STORE PAIN RELIEVER EXTRA STRENGTH

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66715-9737
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
STARCH, CORN (UNII: O8232NY3SJ)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics			
Color	white	Score	no score
Shape	ROUND	Size	12mm
Flavor		Imprint Code	FR33
Contains			

Packaging				
# Item Code	de Package Description	Marketing Start Date	Marketing End Date	
1 NDC:66715-9737-3	3 in 1 CARTON	01/27/2011	08/30/2021	
1	2 in 1 POUCH; Type 0: Not a Combination Product			
NDC:66715- 9737-4	10 in 1 VIAL; Type 0: Not a Combination Product	01/27/2011		
9737-4		01/2//2011		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part343	01/27/2011		

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

Revised: 11/2022 Lil' Drug Store Products, Inc.