

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 caplet)

Acetaminophen 500 mg

Purpose

Pain reliever/fever reducer

Uses

- temporarily relieves minor aches and pains due to:
 - headache
 - the common cold
 - 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 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 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
 - do not take for more than 10 days unless directed by a doctor
- children under 12 years: do not use

Other information

- store between 68-77°F (20-25°C)

Inactive ingredients

corn starch, croscarmellose sodium ¹, hypromellose ¹, lactose monohydrate ¹, magnesium stearate ¹, maltodextrin ¹, medium-chain triglycerides ¹, mineral oil ¹, polydextrose ¹, polyethylene glycol ¹, polyvinyl alcohol ¹, povidone, purified water ¹, sodium starch glycolate ¹, stearic acid ¹, talc ¹, titanium dioxide

1 contains one or more of these ingredients

Questions or comments?

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

Product distributed by:

Lil' Drug Store Products, Inc.
9300 Earhart Lane SW
Cedar Rapids, IA 52404

PRINCIPAL DISPLAY PANEL - 500 mg Caplet Bottle Carton - NDC 66715-6817-4

QUALITY
GUARANTEED

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

Extra Strength

Pain Reliever

Acetaminophen

Pain Reliever/Fever Reducer, 500 mg

ACTUAL SIZE

24
Caplets

Lil'
Drug Store®

Extra Strength Pain Reliever

Acetaminophen

Pain Reliever/Fever Reducer, 500 mg

24
Caplets



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

Extra Strength Pain Reliever

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ACTUAL SIZE

24
Caplets



Product distributed by:
Lil' Drug Store Products, Inc.
9300 Earhart Lane SW
Cedar Rapids, IA 52404

PLD-E108D
FC005264



Lot No:
Exp Date:

KEEP OUTER CARTON FOR COMPLETE
WARNINGS AND PRODUCT INFORMATION.

TAMPER EVIDENT: DO NOT USE IF PRINTED
SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

Drug Facts

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Drug Facts (continued)

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

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QUALITY
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Compare to the
 Active Ingredient in
 Tylenol® Extra Strength**

Extra Strength

Pain Reliever

Acetaminophen

Pain Reliever/Fever Reducer, 500 mg

ACTUAL SIZE

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Lil'

Drug Store[®]

Extra Strength Pain Reliever

Acetaminophen

Pain Reliever/Fever Reducer, 500 mg

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Extra Strength Pain Reliever

Acetaminophen

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PRINCIPAL DISPLAY PANEL - 500 mg Caplet Bottle Carton - NDC 66715-6837-4

QUALITY
 GUARANTEED

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

Extra Strength

Pain Reliever

Acetaminophen

Pain Reliever/Fever Reducer, 500 mg

ACTUAL SIZE

24

Caplets

Lil'

Drug Store[®]

Extra Strength Pain Reliever

Acetaminophen

Pain Reliever/Fever Reducer, 500 mg

24
Caplets



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

Extra Strength Pain Reliever

Acetaminophen

Pain Reliever/Fever Reducer, 500 mg



ACTUAL SIZE

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Caplets



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LIL DRUG STORE PAIN RELIEVER EXTRA STRENGTH

acetaminophen tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66715-6817
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg
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Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
MINERAL OIL (UNII: T5L8T28FGP)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
WATER (UNII: 059QF0KO0R)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white	Score	no score
Shape	OVAL	Size	18mm
Flavor		Imprint Code	P500
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66715-6817-4	1 in 1 CARTON	04/15/2019	08/12/2024
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 not final	part343	01/10/2015	08/12/2024

LIL DRUG STORE PAIN RELIEVER EXTRA STRENGTH

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66715-6827
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
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Inactive Ingredients	
Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
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MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
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MINERAL OIL (UNII: T5L8T28FGP)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
WATER (UNII: 059QF0KO0R)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	white	Score	no score
Shape	OVAL	Size	18mm
Flavor		Imprint Code	AV;0821
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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OTC monograph not final	part343	01/10/2015	08/12/2024

LIL DRUG STORE PAIN RELIEVER EXTRA STRENGTH

acetaminophen tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66715-6837
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
MINERAL OIL (UNII: T5L8T28FGP)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
WATER (UNII: 059QF0KO0R)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white	Score	no score
Shape	OVAL	Size	18mm
Flavor		Imprint Code	TCL;341
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66715-6837-4	1 in 1 CARTON	04/15/2019	08/12/2024
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OTC monograph not final	part343	01/10/2015	08/12/2024

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

Revised: 12/2021

Lil' Drug Store Products, Inc.