

**PAIN RELIEF- acetaminophen tablet, film coated  
DOLGENCORP, LLC**

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**Dollar General 44-531C**

***Active ingredient (in each tablet)***

Acetaminophen 500 mg

***Purpose***

Pain reliever/fever reducer

***Uses***

- temporarily relieves minor aches and pains due to
  - headache
  - toothache
  - muscular aches
  - backache
  - the common cold
  - minor pain of arthritis
  - 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:

- blisters
- rash
- skin reddening

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.**

In case of overdose, get medical help or contact a Poison Control Center right away. Prompt 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**
- 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 take for more than 10 days unless directed by a doctor
- children under 12 years: ask a doctor

***Other information***

- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

***Inactive ingredients***

corn starch, D&C red #27 aluminum lake, D&C yellow #10 aluminum lake, FD&C blue #1 aluminum lake, polyethylene glycol, polyvinyl alcohol, povidone, sodium starch glycolate\*, stearic acid, sucralose, talc, titanium dioxide

\*may contain this ingredient

***Questions or comments?***

**1-888-309-9030**

***Principal display panel***

**DG® | health**

Compare to  
active ingredient of  
Extra Strength  
Tylenol®\*\*

Extra Strength

**Pain Relief**

**Acetaminophen**, 500 mg  
Pain Reliever/Fever Reducer  
• Contains no aspirin

**500**

mg each

**50 Coated Tablets**

Actual Tablet Size

**TAMPER EVIDENT: DO NOT USE IF IMPRINTED  
SAFETY SEAL UNDER CAP IS BROKEN OR MISSING**

\*\*This product is not manufactured or distributed by  
Kenvue Inc., owner of the registered trademark Extra  
Strength Tylenol®.

50844 ORG112353115

DISTRIBUTED BY  
OLD EAST MAIN CO.  
100 MISSION RIDGE  
GOODLETTSVILLE, TN 37072

**100%**

**Satisfaction  
Guaranteed!**

If you're not  
satisfied with  
this product for  
any reason,  
please call us  
so we can  
make you  
100% satisfied.  
(888) 309-9030



Dollar General 44-531C

## PAIN RELIEF

acetaminophen tablet, film coated

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:55910-135
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)		ACETAMINOPHEN	500 mg	
<b>Inactive Ingredients</b>				
Ingredient Name		Strength		
STARCH, CORN (UNII: O8232NY3SJ)				
D&C RED NO. 27 ALUMINUM LAKE (UNII: ZK64F7XSTX)				
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)				
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)				
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
SUCRALOSE (UNII: 96K6UQ3ZD4)				
TALC (UNII: 7SEV7J4R1U)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
<b>Product Characteristics</b>				
Color	red	Score	no score	
Shape	ROUND	Size	11mm	
Flavor		Imprint Code	44;531	
Contains				
<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-135-15	1 in 1 BOX	06/07/2024	
1		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
<b>Marketing Information</b>				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M013	06/07/2024		

**Labeler** - DOLGENCORP, LLC (068331990)

### Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(55910-135)

### Establishment

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		832867837	manufacture(55910-135)

## **Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		832867894	manufacture(55910-135)

## **Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		868734088	manufacture(55910-135)

## **Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		967626305	pack(55910-135)

Revised: 6/2025

DOLGENCORP, LLC