

**PAIN RELIEVER- acetaminophen tablet**  
**Chain Drug Consortium**

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**Premier Value 44-104**

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

Acetaminophen 325 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

- adult takes more than 4,000 mg of acetaminophen in 24 hours
- child takes more than 5 doses in 24 hours
- taken with other drugs containing acetaminophen
- adult has 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 the user has**

liver disease.

**Ask a doctor or pharmacist before use if the user is**

taking the blood thinning drug warfarin.

**Stop use and ask a doctor if**

- pain gets worse or lasts more than 10 days in adults
- new symptoms occur
- pain gets worse or lasts more than 5 days in children under 12 years
- fever gets worse or lasts more than 3 days
- 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 accidental 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	<ul style="list-style-type: none"><li>■ take 2 tablets every 4 to 6 hours while symptoms last</li><li>■ do not take more than 10 tablets in 24 hours</li><li>■ do not take for more than 10 days unless directed by a doctor</li></ul>
children 6-11 years	<ul style="list-style-type: none"><li>■ take 1 tablet every 4 to 6 hours while symptoms last</li><li>■ do not take more than 5 tablets in 24 hours</li><li>■ do not take for more than 5 days unless directed by a doctor</li></ul>
children under 6 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, povidone, sodium starch glycolate, stearic acid

***Questions or comments?***

**1-800-426-9391**

***Principal display panel***

***Premier  
Value®***

**\*COMPARE TO THE ACTIVE INGREDIENT  
IN TYLENOL® REGULAR STRENGTH**

REGULAR STRENGTH

***Pain Reliever***

**ACETAMINOPHEN 325 mg**

**PAIN RELIEVER/FEVER REDUCER**

actual  
size

**100** Tablets-325 mg each

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

\*This product is not manufactured or distributed by  
Johnson & Johnson Corporation, owner of the registered  
trademark Tylenol® Regular Strength.

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**Distributed By:  
Pharmacy Value Alliance, LLC  
407 East Lancaster Avenue,  
Wayne, PA 19087**

If for any reason you are not satisfied with  
this product, please return it to the store  
where purchased for a full refund.



**Premier Value 44-104**

**PAIN RELIEVER**

acetaminophen tablet

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68016-541
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name		Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)		ACETAMINOPHEN	325 mg	
<b>Inactive Ingredients</b>				
Ingredient Name			Strength	
STARCH, CORN (UNII: O8232NY3SJ)				
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)				
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
<b>Product Characteristics</b>				
Color	white	Score	2 pieces	
Shape	ROUND	Size	10mm	
Flavor		Imprint Code	44;104	
Contains				
<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-541-01	1 in 1 CARTON	05/12/2023	
1		100 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	05/12/2023		

**Labeler** - Chain Drug Consortium (101668460)

### Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(68016-541)

### Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(68016-541)

### Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(68016-541)

