

**PAIN RELIEF EXTRA STRENGTH, NON-ASPIRIN- acetaminophen tablet, coated**  
**CHAIN DRUG MARKETING ASSOCIATION INC**

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**Quality Choice 44-519 Delisted**

***Active ingredient (in each gelcap)***

Acetaminophen 500 mg

***Purpose***

Pain reliever/fever reducer

***Uses***

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

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 gels every 6 hours while symptoms last
  - do not take more than 6 gels 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***

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

## ***Inactive ingredients***

croscarmellose sodium, D&C red #33, FD&C blue #1, FD&C red #40, gelatin, hydroxypropyl cellulose, hypromellose, iron oxide black, iron oxide red, iron oxide yellow, polyethylene glycol, povidone, pregelatinized starch, propylene glycol, shellac glaze, stearic acid, titanium dioxide

## ***Questions or comments?***

**1-800-426-9391**

## Principal Display Panel

**QC®  
QUALITY  
CHOICE**

NDC 63868-987-50

**\*Compare to the active  
ingredient in Extra  
Strength TYLENOL®  
Rapid Release Gels**

Extra Strength, Non-Aspirin  
**Pain Relief**  
**Acetaminophen, 500 mg**  
Pain Reliever | Fever Reducer

**50** Gelcaps (500 mg Each)

Actual Size

**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 Extra Strength Tylenol® Rapid Release Gels.  
50844      REV0322E51915

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Novi, MI 48375  
[www.qualitychoice.com](http://www.qualitychoice.com)  
Questions: 800-935-2362

SATISFACTION  
GUARANTEED®  
100%  
**QC**



Quality Choice 44-519

## PAIN RELIEF EXTRA STRENGTH, NON-ASPIRIN

acetaminophen tablet, coated

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-987
Route of Administration	ORAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
D&C RED NO. 33 (UNII: 9DBA05BB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B71O)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

**Product Characteristics**

<b>Color</b>	red, blue	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	19mm
<b>Flavor</b>		<b>Imprint Code</b>	L;5
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-987-50	1 in 1 CARTON	05/10/2004	04/30/2028
1		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:63868-987-10	1 in 1 CARTON	05/10/2004	04/30/2028
2		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
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OTC Monograph Drug	M013	05/10/2004	04/30/2028
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**Labeler -** CHAIN DRUG MARKETING ASSOCIATION INC (011920774)

**Establishment**

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	manufacture(63868-987) , pack(63868-987)

**Establishment**

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(63868-987)

**Establishment**

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(63868-987)

**Establishment**

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(63868-987)

**Establishment**

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(63868-987)

Revised: 1/2026

CHAIN DRUG MARKETING ASSOCIATION INC