

# **PAIN RELIEF EXTRA STRENGTH- acetaminophen tablet, coated H E B**

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**HEB 44-519**

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

Acetaminophen 500 mg

## ***Purpose***

Pain reliever/fever reducer

## ***Uses***

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

## **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
- redness or swelling is present
- any new symptoms appear

**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***

- use by expiration date on package
- 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***

**Compare to Tylenol® Extra Strength Rapid Release Gels** active ingredient\*

NDC 37808-195-20

**H-E-B®**

**TEXAS-SIZE PACK  
225 CT**

**Extra Strength  
Pain Relief**

**Acetaminophen, 500 mg**  
Pain Reliever/Fever Reducer

**225 GELCAPS**

actual  
size

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

**H•E•B®**

**100%**

**GUARANTEE**

***promise***

If you aren't completely pleased  
with this product, we'll be happy to  
replace it or refund your money.  
You have our word on it.

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

50844 REV0525D51920 49903-2510

**MADE WITH PRIDE AND CARE FOR  
H-E-B®, SAN ANTONIO, TX 78204**

Compare to Tylenol® Extra Strength Rapid Release Gels active ingredient\*

ACETAMINOPHEN 500 mg GELCAPS

NDC 37808-195-20



**Extra Strength Pain Relief**  
Acetaminophen, 500 mg

Pain Reliever/Fever Reducer



actual size

**225 GELCAPS**

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

**Drug Facts**

**Active ingredient (in each gelcap)**  
Acetaminophen 500 mg ..... Pain reliever/fever reducer

**Purpose**

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

**100% GUARANTEE promise**  
 If you aren't completely pleased with this product, we'll be happy to replace it or refund your money. You have our word on it.

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

50844 REV0525D51920 49903-2510

MADE WITH PRIDE AND CARE FOR H-E-B®, SAN ANTONIO, TX 78204



IMPRINT AREA  
Lot no/Exp date

PEEL HERE FOR MORE DRUG FACTS

**Drug Facts (continued)**  
 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.  
 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  
 ■ redness or swelling is present  
 ■ any new symptoms appear  
 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 gelcaps every 6 hours while symptoms last  
 ■ do not take more than 6 gelcaps 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)  
 ■ avoid high humidity  
 ■ use by expiration date on package  
**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, stearic acid, stearic acid, titanium dioxide  
**Questions or comments?** 1-800-426-9391

STOP PEELING

HEB 44-519

PAIN RELIEF EXTRA STRENGTH			
acetaminophen tablet, coated			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-195
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
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)			
D&C RED NO. 33 (UNII: 9DBA0SBB0L)			

<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>GELATIN, UNSPECIFIED</b> (UNII: 2G86QN327L)	
<b>HYDROXYPROPYL CELLULOSE, UNSPECIFIED</b> (UNII: 9XZ8H6N6OH)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>FERROSO FERRIC OXIDE</b> (UNII: XM0M87F357)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<b>FERRIC OXIDE YELLOW</b> (UNII: EX438O2MRT)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>SHELLAC</b> (UNII: 46N107B710)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>TITANIUM DIOXIDE</b> (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:37808-195-08	1 in 1 CARTON	05/10/2004	
1		24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:37808-195-12	1 in 1 CARTON	05/10/2004	
2		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:37808-195-20	225 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/10/2004	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	05/10/2004	

**Labeler** - H E B (007924756)

### Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	manufacture(37808-195) , pack(37808-195)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(37808-195)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(37808-195)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(37808-195)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(37808-195)

Revised: 1/2026

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