PAIN RELIEVER- acetaminophen tablet, film coated
L&R Distributors, 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.

Pain Reliever

Active ingredient (in each caplet)

Acetaminophen 500 mg

Purpose

Pain reliever/fever reducer

Uses

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

Warnings

Liver warning: This product contains acetaminophen. The maximum daily dose of this product is 6 caplets (3,000 mg) in 24 hours. 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 accidental overdose, get medical help or contact a Poison Control Center (1-800-222-1222) 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 caplets every 6 hours while symptoms last
  - do not take more than 6 caplets 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 25ºC (77ºF); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive Ingredients

caster oil, hypromellose, povidone, sodium starch glycolate, starch, stearic acid

Questions or comments?

1-800-426-9391

Principal Display Panel

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NDC 15127-735-16

EXTRA STRENGTH/NON-ASPIRIN

PAIN RELIEVER

ACETAMINOPHEN PAIN RELIEVER/FEVER REDUCER

***Compare to the Active Ingredient of Extra Strength Tylenol® Caplets**

50 CAPLETS 500 mg EACH

*This product is not manufactured or distributed by McNeil Consumer Healthcare, owner of the registered trademark Extra Strength Tylenol® Caplets.
50844 REV0715B17515

SATISFACTION GUARANTEED
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SELECT BRAND® DISTRIBUTORS
Pine Bluff, AR 71603 USA
AC (870) 535-3635

Select Brand 44-175

PAIN RELIEVER

acetaminophen tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:15127-735
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**
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POVIDONE (UNII: FZ989GH94E)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
CASTOR OIL (UNII: D5340Y2I9G)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics			
Color	WHITE	**Score**	no score
Shape	OVAL (Tablet)	**Size**	17mm
Flavor		**Imprint Code**	44;175
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:15127-735-19	2 in 1 CARTON	04/02/1993	
1		4 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:15127-735-09	1 in 1 CARTON	04/02/1993	
2		24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:15127-735-16	1 in 1 CARTON	04/02/1993	
3		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:15127-735-08	1 in 1 CARTON	04/02/1993	
4		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
5	NDC:15127-735-05	500 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/02/1993	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part343	04/02/1993	

Labeler - L&R Distributors, Inc. (012578514)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	MANUFACTURE(15127-735)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	PACK(15127-735)

Revised: 5/2017 L&R Distributors, Inc.