

**ACETAMINOPHEN EXTRA STRENGTH PM- acetaminophen, diphenhydramine hcl tablet, film coated  
SAVE-A-LOT FOOD STORES, LTD.**

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**Being Well 44-235**

***Active ingredients (in each caplet)***

Acetaminophen 500 mg  
Diphenhydramine HCl 25 mg

***Purpose***

Pain reliever  
Nighttime sleep-aid

***Uses***

temporary relief of occasional headaches and minor aches and pains with accompanying sleeplessness

***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 product containing diphenhydramine, even one used on skin
- in children under 12 years of age
- 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 have ever had an allergic reaction to this product or any of its ingredients

***Ask a doctor before use if you have***

- a breathing problem such as emphysema or chronic bronchitis
- liver disease
- glaucoma
- difficulty in urination due to enlargement of the prostate gland

***Ask a doctor or pharmacist before use if you are***

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers

***When using this product***

- avoid alcoholic beverages
- drowsiness will occur
- do not drive a motor vehicle or operate machinery

***Stop use and ask a doctor if***

- sleeplessness persists continuously for more than 2 weeks. Insomnia may be a symptom of a serious underlying medical illness.
- new symptoms occur
- redness or swelling is present

- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days

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: take 2 caplets at bedtime. Do not take more than 2 caplets of this product in 24 hours.
- children under 12 years: do not use

**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**

colloidal silicon dioxide, corn starch, croscarmellose sodium, FD&C blue #1 aluminum lake, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, stearic acid, talc, titanium dioxide

**Questions or comments?**

**1-800-426-9391**

**Principal Display Panel**

being  
well®

Compare to **EXTRA STRENGTH  
TYLENOL® PM** active ingredients\*

**EXTRA STRENGTH**

**Acetaminophen PM**

Caplets, 500 mg / Diphenhydramine HCl, 25 mg  
Pain Reliever / Nighttime Sleep-Aid

Non-Habit  
Forming

**50**

Caplets

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® PM.

50844 REV0521G23515

**DISTRIBUTED BY:**

SAVE A LOT LTD.  
ST. ANN, MO 63074

Being Well 44-235

ACETAMINOPHEN EXTRA STRENGTH PM				
acetaminophen, diphenhydramine hcl tablet, film coated				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:46994-935	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)		ACETAMINOPHEN	500 mg	
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)		DIPHENHYDRAMINE HYDROCHLORIDE	25 mg	
Inactive Ingredients				
Ingredient Name			Strength	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
STARCH, CORN (UNII: O8232NY3SJ)				
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)				
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)				
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)				
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
TALC (UNII: 7SEV7J4R1U)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
Product Characteristics				
Color	blue	Score	no score	
Shape	OVAL	Size	17mm	
Flavor		Imprint Code	44;235	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46994-935-15	1 in 1 CARTON	05/15/1994	
1		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M013	05/15/1994	

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(46994-935)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(46994-935) , pack(46994-935)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(46994-935)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(46994-935)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(46994-935)