

**ARTHRITIS PAIN RELIEVER EXTENDED RELEASE- acetaminophen tablet**  
**P & L Development, LLC**

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**Drug Facts**

**Active ingredient (in each caplet)**

**Acetaminophen USP, 650 mg**

**Purpose**

**Pain reliever/fever reducer**

**Uses**

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

**Warnings**

**Liver warning:** This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 6 caplets in 24 hours, which is the maximum daily amount
- 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 a 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.**

**Overdose warning:** In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Quick 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 (see Overdose warning)**
- **adults**
  - take 2 caplets every 8 hours with water
  - swallow whole; do not crush, chew, split or dissolve
  - do not take more than 6 caplets in 24 hours
  - do not use for more than 10 days unless directed by a doctor
- **under 18 years of age:** ask a doctor

## **Other information**

- store between 20-25°C (68-77°F)

## **Inactive ingredients**

hydroxyethyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, pregelatinized starch, sodium starch glycolate, stearic acid

## **Questions or comments?**

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

## **Principal Display Panel**

Compare to the active ingredient in **Tylenol® 8HR Arthritis Pain**†

### **see new warning**

arthritis pain reliever

acetaminophen extended-release tablets USP, 650 mg

pain reliever/fever reducer

- for the temporary relief of minor arthritis pain
- contains no aspirin

caplets\*

(\*capsule-shaped bi-layer tablets)

†This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Tylenol® 8 HR Arthritis Pain.

**TAMPER EVIDENT: DO NOT USE IF FOIL INNER SEAL IS BROKEN OR MISSING.**

**KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.**

Distributed by:

PL Developments

200 Hicks Street

Westbury, NY 11590

## **Product Label**



Compare to the active ingredient in  
**Tylenol® 8 HR Arthritis Pain†**  
 NDC 59726-013-10

**see new warning**  
**arthritis pain reliever**

acetaminophen extended-release tablets  
 USP, 650 mg  
 pain reliever/fever reducer

- for the temporary relief of minor arthritis pain
- contains no aspirin

**100 caplets\***  
 650 mg each (\*capsule-shaped bi-layer tablets)

not actual size



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**Active ingredient (in each caplet)**      **Purpose**  
 Acetaminophen USP, 650 mg.....fever reducer  
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**Uses**

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- if you are allergic to acetaminophen or any of the inactive ingredients in this product

**Drug Facts (continued)**

**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.**

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  - redness or swelling is present
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**KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.**

Exp. Date:  
 Lot No.:

PLD-A683A F0007025



Distributed by:  
**PL Developments**  
 200 Hicks Street  
 Westbury, NY 11590

**READYinCASE Arthritis Pain Reliever**

**ARTHRITIS PAIN RELIEVER EXTENDED RELEASE**

acetaminophen tablet

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:59726-013
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg

**Inactive Ingredients**

Ingredient Name	Strength
<b>HYDROXYETHYL CELLULOSE (140 MPA.S AT 5%)</b> (UNII: 8136Y38GY5)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>SODIUM STARCH GLYCOLATE TYPE A CORN</b> (UNII: AG9B65PV6B)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	

**Product Characteristics**

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	CAPSULE	<b>Size</b>	19mm
<b>Flavor</b>		<b>Imprint Code</b>	G650
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59726-013-10	1 in 1 BOX	04/30/2020	04/30/2026
1		100 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
ANDA	ANDA211544	04/30/2020	04/30/2026

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**Labeler** - P & L Development, LLC (800014821)

Revised: 10/2024

P & L Development, LLC