**PAIN RELIEF EXTRA STRENGTH- acetaminophen tablet**
**P & L Development, LLC**

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**Drug Facts**

**Active ingredient (in each caplet)**

**Acetaminophen 500 mg**

**Purpose**

**Pain reliever/fever reducer**

**Uses**

- temporarily relieves minor aches and pains due to:
  - headache
  - the common cold
  - backache
  - minor pain of arthritis
  - toothache
  - muscular aches
  - 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 ever 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 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 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
  - do not take for more than 10 days unless directed by a doctor
- children under 12 years: do not use

**Other information**

- store between 20-25°C (68-77°F)

**Inactive ingredients**

corn starch, croscarmellose sodium*, hypromellose*, lactose monohydrate*, magnesium stearate*,maltodextrin*, medium-chain triglycerides*, mineral oil*, polydextrose*, polyethylene glycol*, polyvinyl alcohol*, povidone, purified water*, sodium starch glycolate, stearic acid*, talc*, titanium dioxide

*contains one or more of these ingredients

**Questions or comments?**

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

**Principal Display Panel**

Compare to the active ingredient in **Tylenol® Extra Strength**†

Extra Strength

**Pain Reliever**

**Acetaminophen,** 500 mg

Pain Reliever/Fever Reducer

Caplets

†This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Tylenol® Extra Strength.

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

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

**Distributed by:**

**PL Developments**

**200 Hicks Street**

**Westbury, NY 11590**

**Product Label**

**READYinCASE Pain Reliever**

# PAIN RELIEF EXTRA STRENGTH

acetaminophen tablet

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:59726-494 |
| **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** |
| **STARCH, CORN** (UNII: O8232NY3SJ) | |
| **CROSCARMELLOSE SODIUM** (UNII: M28OL1HH48) | |
| **HYPROMELLOSES** (UNII: 3NXW29V3WO) | |
| **LACTOSE MONOHYDRATE** (UNII: EWQ57Q8I5X) | |
| **MAGNESIUM STEARATE** (UNII: 70097M6I30) | |
| **MINERAL OIL** (UNII: T5L8T28FGP) | |
| **POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WQ0SDW1A) | |
| **POVIDONE** (UNII: FZ989GH94E) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **SODIUM STARCH GLYCOLATE TYPE A CORN** (UNII: AG9B65PV6B) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |
| **TALC** (UNII: 7SEV7J4R1U) | |
| **TITANIUM DIOXIDE** (UNII: 15FIX9V2JP) | |
| **POLYVINYL ALCOHOL, UNSPECIFIED** (UNII: 532B59J990) | |
| **MALTODEXTRIN** (UNII: 7CVR7L4A2D) | |
| **MEDIUM-CHAIN TRIGLYCERIDES** (UNII: C9H2L21V7U) | |
| **POLYDEXTROSE** (UNII: VH2XOU12IE) | |

| Product Characteristics | | | |
|---|---|---|---|
| **Color** | white | **Score** | no score |
| **Shape** | CAPSULE | **Size** | 18mm |
| **Flavor** | | **Imprint Code** | TCL341;AV;0821;P500 |
| **Contains** | | | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:59726-494-24 | 1 in 1 BOX | 03/31/2016 | 12/31/2026 |
| | | 24 in 1 BOTTLE, PLASTIC; Type 0: Not a | | |

| | | | | |
|---|---|---|---|---|
| **1** | | 24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | | |
| **2** | NDC:59726-494-05 | 1 in 1 BOX | 03/31/2016 | 12/31/2026 |
| **2** | | 500 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | | |
| **3** | NDC:59726-494-10 | 100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 03/31/2016 | 12/31/2026 |
| **4** | NDC:59726-494-50 | 1 in 1 BOX | 03/31/2016 | 12/31/2026 |
| **4** | | 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 | 03/31/2016 | 12/31/2026 |

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

Revised: 8/2025 P & L Development, LLC