

**ACETAMINOPHEN- acetaminophen tablet, film coated**  
**ARMY AND AIR FORCE EXCHANGE SERVICE**

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**Pain Reliever**

***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
  - toothache
  - backache
  - muscular aches
  - minor pain of arthritis
  - 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**

- if you are allergic to acetaminophen or any of the inactive ingredients in this product
- 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
- 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 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 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

***Inactive ingredients***

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

***Questions or comments?***

**1-800-426-9391**

**Principal Display Panel**

**exchange✓ select™**

Compare To The Active Ingredient of  
Extra Strength Tylenol® Caplets†

Contains no aspirin

**ACETAMINOPHEN  
EXTRA  
STRENGTH**

*Pain Reliever  
Fever Reducer*

**50 CAPLETS  
500 mg each**

Actual Size

✓ quality  
value

**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® Caplets.

**"SATISFACTION GUARANTEED OR YOUR MONEY BACK"**

Manufactured For Your Military Exchanges  
Distributed by: LNK International, Inc.,  
Hauppauge, NY 11788  
1-800-426-9391

50844      REV0621P17515



Exchange Select 44-175

## ACETAMINOPHEN

acetaminophen tablet, film coated

### Product Information

|                         |                |                    |               |
|-------------------------|----------------|--------------------|---------------|
| Product Type            | HUMAN OTC DRUG | Item Code (Source) | NDC:55301-175 |
| 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 |
|--|----------|
| CASTOR OIL (UNII: D5340Y2I9G)                            |          |
| HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)             |          |
| POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)                 |          |
| SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2) |          |
| STARCH, CORN (UNII: O8232NY3SJ)                          |          |
| STEARIC ACID (UNII: 4ELV7Z65AP)                          |          |

Product Characteristics

|          |       |              |          |
|----------|-------|--------------|----------|
| Color    | white | Score        | no score |
| Shape    | OVAL  | Size         | 17mm     |
| Flavor   |       | Imprint Code | 44;175   |
| Contains |       |              |          |

Packaging

| # | Item Code        | Package Description   | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:55301-175-95 | 10 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product  | 04/02/1993           |                    |
| 2 | NDC:55301-175-15 | 1 in 1 CARTON   | 04/02/1993           |                    |
| 2 |                  | 50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product  |                      |                    |
| 3 | NDC:55301-175-60 | 200 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 04/02/1993           |                    |
| 4 | NDC:55301-175-03 | 1 in 1 CARTON   | 04/02/1993           | 01/17/2024         |
| 4 |                  | 10 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product  |                      |                    |
| 5 | NDC:55301-175-12 | 1 in 1 CARTON   | 04/02/1993           | 01/26/2017         |
| 5 |                  | 100 in 1 BOTTLE; Type 0: Not a Combination Product          |                      |                    |
| 6 | NDC:55301-175-06 | 2 in 1 CARTON   | 04/02/1993           | 01/26/2017         |
| 6 |                  | 100 in 1 BOTTLE; 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                                     | 04/02/1993           |                    |

**Labeler** - ARMY AND AIR FORCE EXCHANGE SERVICE (001695568)

## Establishment

| Name                    | Address | ID/FEI    | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. |         | 038154464 | pack(55301-175)     |

## Establishment

| Name                    | Address | ID/FEI    | Business Operations    |
|-------------------------|---------|-----------|------------------------|
| LNK International, Inc. |         | 832867837 | manufacture(55301-175) |

## Establishment

| Name                    | Address | ID/FEI    | Business Operations    |
|-------------------------|---------|-----------|------------------------|
| LNK International, Inc. |         | 832867894 | manufacture(55301-175) |

## Establishment

| Name                    | Address | ID/FEI    | Business Operations    |
|-------------------------|---------|-----------|------------------------|
| LNK International, Inc. |         | 868734088 | manufacture(55301-175) |

## Establishment

| Name                    | Address | ID/FEI    | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. |         | 967626305 | pack(55301-175)     |

## Establishment

| Name                    | Address | ID/FEI    | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. |         | 117597853 | pack(55301-175)     |

Revised: 1/2024

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