**FIRST AID ONLY EXTRA-STRENGTH NON-ASPIRIN- acetaminophen tablet, film coated**
**Acme United Corporation**

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**First Aid Only Extra-Strength Non-Aspirin**

***Drug Facts***

***Active ingredient (in each tablet)***

Acetaminophen 500 mg

***Purpose***

Pain reliever/fever reducer

***Uses***

**For the temporary relief of minor aches and pains associated with**

- headache
- toothache
- minor arthritis pain
- muscular aches
- common cold
- menstrual cramps

temporarily reduces fever

***Warnings***

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

- more than 8 tablets 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 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.
- for more than 10 days for pain unless directed by a doctor
- for more than 3 days for fever unless directed by a doctor

**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 using and ask a doctor if**
- symptoms do not improve
- pain or fever persists or gets worse
- new symptoms occur
- redness or swelling is present

**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 use more than directed**

**Adults and children: (12 years and older)**

Take 2 tablets every 4 to 6 hours as needed. Do not take more than 8 tablets in 24 hours.

**Children under 12 years:**

Do not give this adult strength product to children under 12 years of age;

this will provide more than the recommended dose (overdose) and may cause liver damage.

***Other information***
- store at room temperature 59º-86ºF (15º-30ºC)
- tamper-evident sealed packets
- do not use any opened or torn packets

***Inactive ingredients***

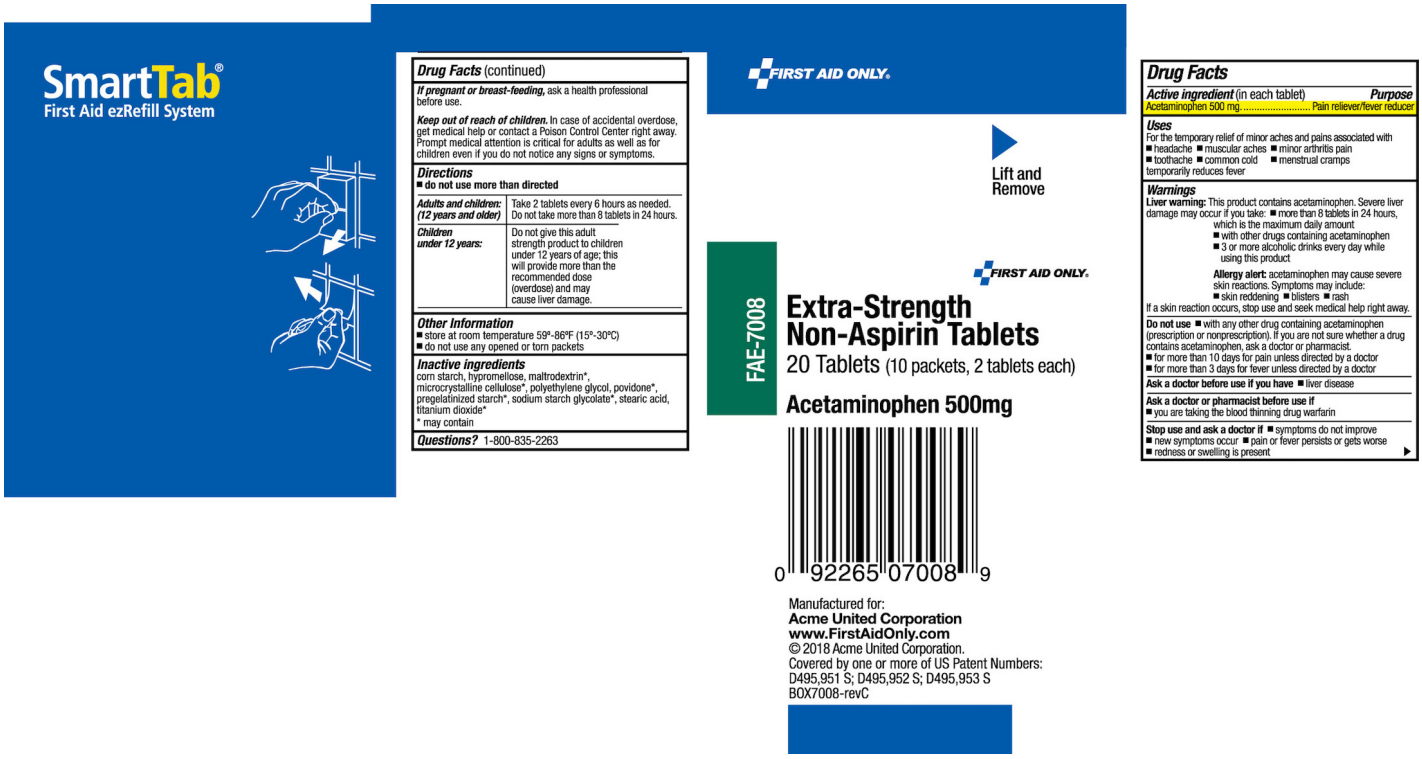
corn starch, hypromellose, maltodextrin*, microcrystalline cellulose*, polyethylene glycol, povidone*, pregelatinized corn starch*, sodium starch glycolate*, stearic acid, titanium dioxide*

* *May contain*

***Questions?*** 1-800-835-2263

## First Aid Only Extra-Strength Non-Aspirin Label

First Aid Only Extra-Strength Non-Aspirin Label




# FIRST AID ONLY EXTRA-STRENGTH NON-ASPIRIN

acetaminophen tablet, film coated

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:0924-0223(NDC:47682-175) |
| **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** |
| **POLYETHYLENE GLYCOL** (UNII: 3WQ0SDW1A) | |
| **HYPROMELLOSES** (UNII: 3NXW29V3WO) | |
| **POVIDONE** (UNII: FZ989GH94E) | |
| **SODIUM STARCH GLYCOLATE TYPE A POTATO** (UNII: 5856J3G2A2) | |

| | |
|---|---|
| **STARCH, CORN** (UNII: O8232NY3SJ) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |
| **TITANIUM DIOXIDE** (UNII: 15FIX9V2JP) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | white (white) | **Score** | no score |
| **Shape** | ROUND (ROUND) | **Size** | 12mm |
| **Flavor** | | **Imprint Code** | AZ;235 |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:0924-0223-01 | 20 in 1 BOX, UNIT-DOSE | 12/30/2008 | |
| **1** | | 2 in 1 PACKET; 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 | 12/30/2008 | |

## Labeler - Acme United Corporation (001180207)

## Registrant - Acme United Corporation (001180207)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Acme United Corporation | | 045924339 | relabel(0924-0223) , repack(0924-0223) |

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Acme United Corporation | | 080119599 | relabel(0924-0223) , repack(0924-0223) |

Revised: 11/2024 Acme United Corporation