**ULINE ACETAMINOPHEN EXTRA STRENGTH- acetaminophen tablet, film coated**
**Uline**

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**Uline Acetaminophen Extra Strength**

## *Drug Facts*

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

Acetaminophen 500 mg

### *Purpose*

Pain reliever/fever reducer

### *Uses*

Temporarily relieves minor aches and pains due to:

- headache
- muscular aches
- minor arthritis pain
- backache
- the common cold
- toothache
- 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 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.

- 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 have**

- liver disease

**Stop use and ask a doctor if**

- pain gets worse or lasts for 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 (1-800-222-1222). 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 use more than directed (see overdose warning)**

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

Take 2 tablets every 6 while symptoms last. Do not take more than 6 tablets in 24 hours, unles directed by a doctor. Do not use for more than 10 days unless directed by a doctor.

**Children under 12 years:**

Ask a doctor.

***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*, polyethylene glycol*, povidone (K-30)*, pregelatinized starch*, purified water*, sodium starch glycolate, stearic acid, titanium dioxide*

* *May contain*

### *Questions or comments?*

1-800-295-5510

### Uline Acetaminophen Extra Strength Label

ULINE

Tamper evident sealed packets:

Do not use if packet is open or torn.

This package is for households without young children

Acetaminophen

Pull to Open

Extra Strength

500 mg

• Pain Reliever

• Fever Reducer

50 Packets

2 tablets Each


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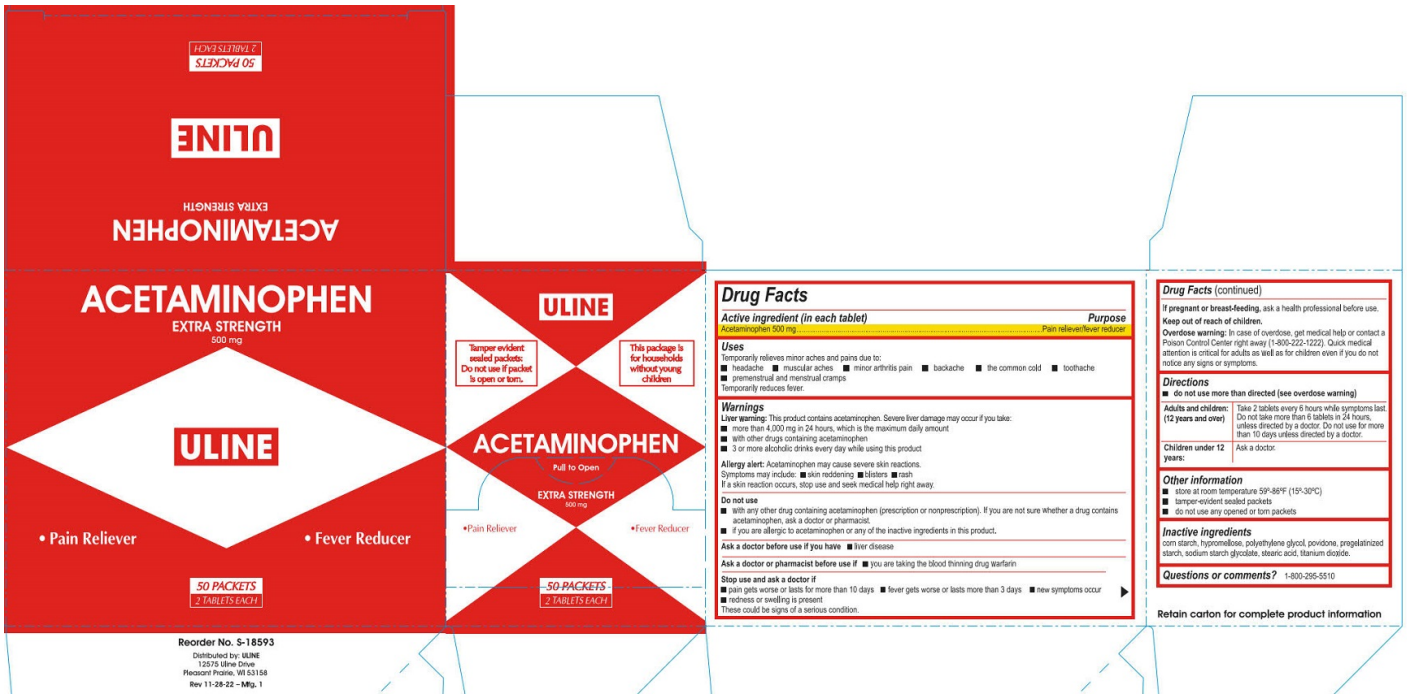

# ULINE ACETAMINOPHEN EXTRA STRENGTH

acetaminophen tablet, film coated

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:69790-173 |
| **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** |
| **POVIDONE** (UNII: FZ989GH94E) | |
| **STARCH, CORN** (UNII: O8232NY3SJ) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |
| **SODIUM STARCH GLYCOLATE TYPE A POTATO** (UNII: 5856J3G2A2) | |

| Product Characteristics | | | |
|---|---|---|---|
| **Color** | white | **Score** | 2 pieces |
| **Shape** | ROUND | **Size** | 12mm |
| **Flavor** | | **Imprint Code** | 44;148 |
| **Contains** | | | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:69790-173-33 | 50 in 1 BOX | 01/30/2024 | |
| **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 | 01/30/2024 | |


# ULINE ACETAMINOPHEN EXTRA STRENGTH

acetaminophen tablet, film coated

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:69790-804 |
| **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** |
| **HYPROMELLOSES** (UNII: 3NXW29V3WO) | |
| **POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WQ0SDW1A) | |
| **SODIUM STARCH GLYCOLATE TYPE A POTATO** (UNII: 5856J3G2A2) | |
| **STARCH, CORN** (UNII: O8232NY3SJ) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |
| **POVIDONE** (UNII: FZ989GH94E) | |
| **TITANIUM DIOXIDE** (UNII: 15FIX9V2JP) | |

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

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:69790-804-33 | 50 in 1 BOX | 10/07/2019 | |
| **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 | 10/07/2019 | |

# ULINE ACETAMINOPHEN EXTRA STRENGTH

acetaminophen tablet, film coated

| Product Information | | | |
|---|---|---|---|

| | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:69790-126 |
| **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, UNSPECIFIED** (UNII: 3WQ0SDW1A) | |
| **SODIUM STARCH GLYCOLATE TYPE A POTATO** (UNII: 5856J3G2A2) | |
| **STARCH, CORN** (UNII: O8232NY3SJ) | |
| **MALTODEXTRIN** (UNII: 7CVR7L4A2D) | |
| **CELLULOSE, MICROCRYSTALLINE** (UNII: OP1R32D61U) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |
| **POVIDONE** (UNII: FZ989GH94E) | |
| **HYPROMELLOSES** (UNII: 3NXW29V3WO) | |

| Product Characteristics | | | |
|---|---|---|---|
| **Color** | white | **Score** | no score |
| **Shape** | ROUND | **Size** | 12mm |
| **Flavor** | | **Imprint Code** | FR;33 |
| **Contains** | | | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:69790-126-33 | 50 in 1 BOX | 10/07/2019 | 11/01/2023 |
| **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 | 10/07/2019 | 11/01/2023 |

# ULINE ACETAMINOPHEN EXTRA STRENGTH

acetaminophen tablet, film coated

## Product Information

| | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:69790-154 |

| | |
|---|---|
| **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) | |
| **SODIUM STARCH GLYCOLATE TYPE A POTATO** (UNII: 5856J3G2A2) | |
| **POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WQ0SDW1A) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |
| **POVIDONE K30** (UNII: U725QWY32X) | |
| **HYPROMELLOSES** (UNII: 3NXW29V3WO) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | white ((White to Off-White)) | **Score** | no score |
| **Shape** | ROUND | **Size** | 12mm |
| **Flavor** | | **Imprint Code** | G552 |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:69790-154-33 | 50 in 1 BOX | 01/30/2024 | |
| **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 | 01/30/2024 | |

## Labeler - Uline (039612668)

## Registrant - Unifirst First Aid Corporation (832947092)

Revised: 9/2025 Uline