

**MAPAP EXTRA STRENGTH- acetaminophen capsule**  
**Major Pharmaceuticals**

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**0617 - Major**

***Drug Facts***

***Active Ingredient (in each capsule)***

Acetaminophen 500 mg

***Purpose***

***Purpose***

Pain reliever/ Fever reducer

***Uses***

Temporarily reduces fever and relieves minor aches and pains due to:

- headache
- muscular aches
- the common cold
- toothache
- backache
- minor pain of arthritis
- premenstrual and menstrual cramps

***Warnings***

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

- more than 6 capsules 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 drugs 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 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 the 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 take more than directed (see overdose warning).**

**adults and children 12 years and over:**

- take 2 capsules every 6 hours while symptoms last
- Do not take more than 6 capsules 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 in a dry place at 15° - 30°C (59° - 86°F).

***Inactive Ingredients***

FD&C Blue #1, FD&C Red #40, Gelatin, Polyvinylpyrrolidone, Pregelatinized Starch, Sodium Starch Glycolate, Stearic Acid and Titanium Dioxide

**Questions?**

**Call (800) 231-4670**

Distributed by: **Major® Pharmaceuticals**

Indianapolis, IN 46268

(800) 616-2471

[www.majorpharmaceuticals.com](http://www.majorpharmaceuticals.com)

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

**Major**

**NDC 0904-1987-60**

**EXTRA STRENGTH**

**Mapap**

**Acetaminophen 500 mg Each**

**CAPSULES SAFETY SEALED FOR YOUR PROTECTION**

**SEE NEW WARNINGS INFORMATION AND DIRECTIONS**

**ASPIRIN FREE**

**Strong Pain & Fever Relief**

**100 CAPSULES**

NDC 0904-1987-60

**MAJOR**<sup>®</sup>

**EXTRA STRENGTH**

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**ASPIRIN FREE**  
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**Drug Facts**


**Active ingredient (in each capsule)** Purpose  
Acetaminophen 500 mg.....Pain reliever/Fever reducer

**Uses**  
Temporarily reduces fever and relieves minor aches and pains due to: • headache • muscular aches • the common cold • toothache • backache • minor pain of arthritis • premenstrual and menstrual cramps

**Warnings**  
**Liver warning:** This product contains acetaminophen. Severe liver damage may occur if you take • more than 6 capsules 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

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Indianapolis, IN 46268  
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**09041 98760 9**  
Lot # Exp. Date

M-29  
Rev. 10/23 3  
RS2053  
Re-Order  
No. 700691



**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

**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 the 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 take more than directed (see overdose warning)
- take 2 capsules every 6 hours while adults and children 12 years and over:
- do not take more than 6 capsules 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 in a dry place at 15° – 30°C (59° – 86°F).

**Inactive ingredients** FD&C blue #1, FD&C red #40, gelatin, polyvinylpyrrolidone, pregelatinized starch, sodium starch glycolate, stearic acid and titanium dioxide

**Questions or comments?**  
Call (800) 231-4670

| <b>MAPAP EXTRA STRENGTH</b>  |                |                           |                 |
|--|----------------|---------------------------|-----------------|
| acetaminophen capsule  |                |                           |                 |
| Product Information  |                |                           |                 |
| <b>Product Type</b>  | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:0904-1987   |
| <b>Route of Administration</b>                                     | ORAL           |                           |                 |
| Active Ingredient/Active Moiety                                    |                |                           |                 |
| <b>Ingredient Name</b>   |                | <b>Basis of Strength</b>  | <b>Strength</b> |
| ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D) |                | ACETAMINOPHEN             | 500 mg          |
| Inactive Ingredients   |                |                           |                 |
| <b>Ingredient Name</b>   |                | <b>Strength</b>           |                 |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD)                                 |                |                           |                 |
| FD&C RED NO. 40 (UNII: WZB9127XOA)                                 |                |                           |                 |
| GELATIN (UNII: 2G86QN327L)   |                |                           |                 |
| POVIDONE K30 (UNII: U725QWY32X)                                    |                |                           |                 |
| STARCH, CORN (UNII: O8232NY35J)                                    |                |                           |                 |

**SODIUM STARCH GLYCOLATE TYPE A POTATO** (UNII: 5856J3G2A2)

**STEARIC ACID** (UNII: 4ELV7Z65AP)

**TITANIUM DIOXIDE** (UNII: 15FIX9V2JP)

### Product Characteristics

|                 |            |                     |          |
|-----------------|------------|---------------------|----------|
| <b>Color</b>    | white, red | <b>Score</b>        | no score |
| <b>Shape</b>    | CAPSULE    | <b>Size</b>         | 22mm     |
| <b>Flavor</b>   |            | <b>Imprint Code</b> | CPC;617  |
| <b>Contains</b> |            |                     |          |

### Packaging

| # | Item Code        | Package Description                                 | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:0904-1987-60 | 100 in 1 BOTTLE; Type 0: Not a Combination Product  | 12/22/1997           |                    |
| 2 | NDC:0904-1987-80 | 1000 in 1 BOTTLE; Type 0: Not a Combination Product | 01/29/2001           | 04/30/2017         |

### Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M013                                     | 12/22/1997           |                    |

**Labeler** - Major Pharmaceuticals (191427277)

Revised: 5/2024

Major Pharmaceuticals