

# **MAPAP ARTHRITIS PAIN- acetaminophen tablet, film coated, extended release**

**Major Pharmaceuticals**

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## **Major Pharmaceuticals Mapap® Arthritis Pain Drug Facts**

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

Acetaminophen 650 mg

### **Purpose**

Pain reliever/fever reducer

### **Uses**

- temporarily relieves minor aches and pains due to:
- minor pain of arthritis
- muscular aches
- backache
- premenstrual and menstrual cramps
- the common cold
- headache
- toothache
- temporarily reduces fever

### **Warnings**

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

- more than 6 caplets 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 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 (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	<ul style="list-style-type: none"><li>• take 2 caplets every 8 hours with water</li><li>• swallow whole; do not crush, chew, split or dissolve</li><li>• do not take more than 6 caplets in 24 hours</li><li>• do not use for more than 10 days unless directed by a doctor</li></ul>
under 18 years of age	<ul style="list-style-type: none"><li>• ask a doctor</li></ul>

**Other information**

- store at 20-25°C (68-77°F)
- **do not use if printed foil under cap is broken or missing**

- meets the requirements of USP *Dissolution Test 2*

### **Inactive ingredients**

carnauba wax, colloidal silicon dioxide, croscarmellose sodium, hypromellose, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, polysorbate 80, povidone, pregelatinized starch, stearic acid, titanium dioxide

### **Questions or comments?**

**1-800-616-2471**

### **Principal Display Panel**

MAJOR<sup>®</sup>

Push & Turn Cap

Mapap<sup>®</sup> ARTHRITIS PAIN

ACTUAL SIZE

ACETAMINOPHEN EXTENDED-RELEASE TABLETS, 650 mg

CAPLETS

PAIN RELIEVER/FEVER REDUCER

FOR THE TEMPORARY RELIEF OF MINOR ARTHRITIS PAIN

Compare to the active ingredient in Tylenol<sup>®</sup> 8HR Arthritis Pain

DO NOT USE WITH OTHER MEDICINES CONTAINING ACETAMINOPHEN

100 Caplets\* 650 mg. Each

\*Capsule-Shaped Tablets

MAJOR<sup>®</sup>

NDC 0904-5769-60

Push & Turn Cap

# Māpap<sup>®</sup>

ARTHRITIS PAIN

ACETAMINOPHEN  
EXTENDED-RELEASE TABLETS, 650 mg  
PAIN RELIEVER/FEVER REDUCER  
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Contains No Aspirin U.S. Patent 7,897,172

Distributed by  
**MAJOR®**  
**PHARMACEUTICALS**  
17177 N Laurel Park Drive  
Suite 233  
Livonia, MI 48152  
M-05 REV. 09/19  
Re-Order No. 100381

**Drug Facts**

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**Drug Facts (continued)**

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**Questions or comments? 1-800-616-2471**

\*\*This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., distributor of Tylenol® 8HR Arthritis Pain.

**Gluten Free**



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**CODE AREA**

54478 5C C10

**MAPAP ARTHRITIS PAIN**

acetaminophen tablet, film coated, extended release

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0904-5769
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg

**Inactive Ingredients**

Ingredient Name	Strength
<b>CARNAUBA WAX</b> (UNII: R12CBM0EIZ)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

**Product Characteristics**

<b>Color</b>	WHITE	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	19mm
<b>Flavor</b>		<b>Imprint Code</b>	L544
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904-5769-60	1 in 1 CARTON	02/16/2006	
1		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
ANDA	ANDA075077	02/16/2006	

**Labeler** - Major Pharmaceuticals (191427277)

Revised: 11/2022

Major Pharmaceuticals