

**ACETAMINOPHEN PAIN RELIEVER, FEVER REDUCER 500 MG- acetaminophen tablet**  
**Medsouce Pharmaceuticals**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**Acetaminophen 500 mg Tablet**

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

Acetaminophen 500 mg

***Purposes***

Pain reliever/fever reducer

***Uses***

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

***Warnings***

**Liver warning:** This product contains acetaminophen. The maximum daily dose of this product is 6 tablets (3,000 mg) in 24 hours. 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

**Do not use**

- with any other drug containing acetaminophen (prescription or non prescription). 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:** Taking more than the recommended dose (overdose) may cause liver damage. 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 tablets every 6 hours while symptoms last
- do not take more than 6 tablets 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

- **Do not use if imprinted safety seal under cap is broken or missing**
- Store at room temperature

### Inactive ingredients

Povidone, Pregelatinized Starch, Sodium Starch Glycolate, Stearic Acid.

### Questions?

If you have any questions or comments, or to report an adverse event, please contact (800) 795-9775.

### Principal Display Panel

**ACETAMINOPHEN TABLETS 500MG**  
 GENERIC FOR TYLENOL ES

LOT: NDC 45865-0136-30 EXP: #30

DO NOT TAKE MORE THAN 6 TABLETS IN 24 HOURS  
 TAKE 1-2 TABLETS EVERY 4-6 HOURS, AS NEEDED FOR PAIN

ACETAMINOPHEN TABLETS 500MG  
 GENERIC FOR TYLENOL ES # 30  
 lot# exp: NDC: 45865-0136-30 MFR NDC: 51645-0706-10

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PATIENT LOG CHART CLAMP

PEEL HERE

# ACETAMINOPHEN PAIN RELIEVER,FEVER REDUCER 500 MG

acetaminophen tablet

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:45865-136(NDC:51645-706)
<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>
POVIDONE (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

## Product Characteristics

<b>Color</b>	white	<b>Score</b>	2 pieces
<b>Shape</b>	ROUND (round flat faced beveled edge)	<b>Size</b>	12mm
<b>Flavor</b>		<b>Imprint Code</b>	GPI;A5
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:45865-136-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2020	

## Marketing Information

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC monograph not final	part343	10/01/2020	

**Labeler** - Medsource Pharmaceuticals (833685915)

## Establishment

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
Medsource Pharmaceuticals		833685915	repack(45865-136)

