

**QAULITY CHOICE PAIN RELIEF EXTRA STRENGTH- acetaminophen liquid**  
**CHAIN DRUG MARKETING ASSOCIATION INC**

*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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**☐Active Ingredient**

☐Acetaminophen 500 mg

**Purpose**

Pain Reliever/Fever Reducer

**☐Uses**

- ☐temporarily relieves minor aches and pains
- temporarily reduces fever

**Warnings**

**Liver warning:** This product contains acetaminophen. The maximum daily dose of this product is 6 tablespoonfuls in 24 hours. Severe liver damage may occur if you take

- more than 8 tablespoonfuls (4,000 mg of acetaminophen) in 24 hours
- 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 before use** if you are taking the blood thinning drug warfarin.

**When using this product:** Do not exceed recommended dose.

**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.** ☐In case of overdose, get medical help or contact a Poison Control Center right away at 1-800-222-1222. Quick medical attention is critical for adults/children even if you do not notice any signs or symptoms.

**Directions Do not take more than directed**

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Age	Dose
Adults and children 12 years of age and older	Take 1 to 2 tablespoonfuls every 4 to 6 hours as needed. not more than 6 tablespoonfuls in 24 hours
Children under 12 years	Do not use

**Inactive ingredients:** artificial and natural cherry flavor, citric acid, FD&C Red #40, methylparaben, polyethylene glycol, propylene glycol, propylparaben, purified water, sodium citrate, sucralose.

**Question or comments? 1-800-540-3765**

**Drug Facts**

**Active ingredient**  
(in each 15mL Tablespoonful)  
Acetaminophen 500 mg.....Pain Reliever / Fever Reducer

**Purposes**

**Uses**

- temporarily relieves minor aches and pains
- temporarily reduces fever

**Warnings**

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- more than 8 tablespoonfuls (4,000 mg of acetaminophen) in 24 hours
- 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 before use if you are taking the blood thinning drug warfarin.**

**When using this product: Do not exceed recommended dose.**

**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.

**QC**  
QUALITY CHOICE

NDC 63868-564-08

Compare to the active ingredient in Extra Strength **TYLENOL**®

**PAIN RELIEF**

**Extra Strength**

**ACETAMINOPHEN**

500 mg / 15 mL

**PAIN RELIEVER • FEVER REDUCER**

- Aspirin free
- Alcohol free

**Cherry Flavor**

**8 FL OZ (237 mL)**

**Drug Facts (continued)**

If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away at 1-800-222-1222. Quick medical attention is critical for adults/children even if you do not notice any signs or symptoms.

**Directions • do not take more than directed**

Age	Dose
Adults and children 12 years of age and older	Take 1 to 2 Tablespoonfuls every 4 to 6 hours as needed; not more than 6 Tablespoonfuls in 24 hours
Children under 12 years	Do not use

**Other information**

- store at room temperature 15°- 30°C (59°-86°F)
- protect from freezing • protect from light
- TAMPER-EVIDENT:** Do not use if foil seal over bottle opening is torn, broken or missing.

**Inactive ingredients:** artificial and natural cherry flavor, citric acid, FD&C Red #40, methylparaben, polyethylene glycol, propylene glycol, propylparaben, purified water, sodium citrate, sucralose

**Questions or comments? 1-800-540-3765**

\*This product is not manufactured or distributed by Johnson & Johnson, owner of the registered trademark Tylenol®.

Satisfaction Guaranteed

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www.qualitychoice.com  
Questions: 800-935-2362

Code # L-148  
Lot#  
Exp. Date:

Rev.:05/20

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## QAULITY CHOICE PAIN RELIEF EXTRA STRENGTH

acetaminophen liquid

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-564
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg in 15 mL

### Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

PROPYLPARABEN (UNII: Z8IX2SC1OH)				
WATER (UNII: 059QF0KO0R)				
SODIUM CITRATE (UNII: 1Q73Q2JULR)				
SUCRALOSE (UNII: 96K6UQ3ZD4)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-564-08	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2020	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final		part343	12/01/2020	

**Labeler** - CHAIN DRUG MARKET ING ASSOCIATION INC (011920774)

Revised: 12/2020

CHAIN DRUG MARKETING ASSOCIATION INC