

PURGENLABS ACETAMINOPHEN- actaminophen liquid
Advanced Rx LLC

Active Ingredient

in each 15 mL Tablespoonful

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.

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

Questions or comments? call toll free 1-800-630-8895

Drug Facts

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Compare to the active ingredient in Tylenol® Extra Strength™

Adult Extra Strength

ACETAMINOPHEN



500 mg per 15 ml

- ✓ Pain Reliever
- ✓ Fever Reducer
- ✓ Alcohol Free

RAPID BURST LIQUID

For ages 12 years and over

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. 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 20°- 25°C (68°-77°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? call toll-free 1-800-630-8895

Manufactured for: ADVANCED RX LLC
1942 NE 163rd St. North Miami Beach, FL 33162 U.S.A

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

Code #: L-230
Rev. 04/24

Lot#
Exp. Date:



PURGENLABS ACETAMINOPHEN

actaminophen liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80513-230
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: WZ B9127XOA)	
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:80513-230-08	1 in 1 CARTON	06/01/2024	
1		237 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	06/01/2024	

Labeler - Advanced Rx LLC (042795108)

Revised: 6/2024

Advanced Rx LLC