EXTRA STRENGTH PAIN RELIEF- acetaminophen suspension Preferred Pharmaceuticals Inc.

GC ES apap liquid

Active ingredient (in each 15mL 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 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

When using this product: Do not exceed recommended dose.

Stop use and ask a doctor if

new symptoms occur

- redness or swelling is present
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days

These could be signs of a serious condition.

If pregnant or breastfeeding, 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. 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

Adults and children Take 1-2 Tablespoons
12 years of age and every 4 to 6 hours as older needed; not more than 6 Tablespoons in 24 hours

Children under 12 Do not use years

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

Relabeled By: Preferred Pharmaceuticals Inc.

PRINCIPAL DISPLAY PANEL

Extra Strength APAP Liquid

500mg per 15mL Genric for Tylenol Liquid

In each 15mL: Acetaminophen 500mg...pain reliever/fever reducer

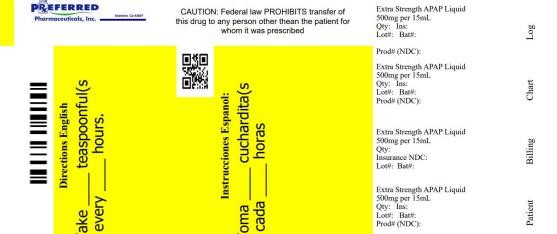
> Pkg Size: Exp Date: Lot#:

Batch#:

Ins: Mfg: Geri-Care Pharmaceutical Corp

Prod#:

FIOGH:
Wming
SEF NEW DOSNG INFORMATION. This product
contains acctaminophen. Severe liver damage may occur
if you take: more than 4,000mg in 24 hours, with other
drugs containing acctaminophen. Do not use with any
other product drug containing acctaminophen (prescripti
acctaminophen or any of the inactive ingredients in this
product. See bottle for complete list of drug facts and
Aspirint Flavor. Cherry Flavor. Siore at room temperature
15°-30°C (59°-86°F). Protect from light. Protect from
freezing.



EXTRA STRENGTH PAIN RELIEF

ORAL

acetaminophen suspension

Product Information

Route of Administration

Product Type HUMAN OTC DRUG NDC:68788-8375(NDC:57896-206) Item Code (Source)

Active Ingredient/Active Moiety

SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2|ULR)

Ingredient Name Basis of Strength Strength ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN 500 mg in 15 mL

Inactive Ingredients Ingredient Name Strength CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP) FD&C RED NO. 40 (UNII: WZB9127XOA) **METHYLPARABEN** (UNII: A2I8C7HI9T) POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) SUCRALOSE (UNII: 96K6UQ3ZD4) PROPYLENE GLYCOL (UNII: 6DC9Q167V3) PROPYLPARABEN (UNII: Z8IX2SC1OH) WATER (UNII: 059QF0KO0R)

Product Characteristics					
Color	red	Score			
Shape		Size			
Flavor		Imprint Code			

Contains

l	Packaging						
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1	NDC:68788- 8375-2	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/24/2023			

Marketing Information						
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date			
OTC Monograph Drug	343	02/24/2023				

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment							
Name	Address	ID/FEI	Business Operations				
Preferred Pharmaceuticals Inc.		791119022	RELABEL(68788-8375)				

Revised: 5/2025 Preferred Pharmaceuticals Inc.