

EXTRA STRENGTH ACETAMINOPHEN- acetaminophen tablet
Preferred Pharmaceuticals Inc.

gc222

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.

Severe liver damage may occur if you take:

- more than 8 caplets (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 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 symptom 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. 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**
- adults and children 12 years and over: take 2 caplets every 6 hours, as needed. Do not take more than 6 caplets in 24 hours. Do not take for more than 10 days unless directed by a doctor.
- children under 12 years: ask a doctor

Other Information

- store at 20°C-25°C (68°F-77°F)
- Repackaged By: Preferred Pharmaceuticals Inc.

Inactive ingredients

corn starch, povidone, pregelatinized starch, sodium starch glycolate, stearic acid

Questions or comments?

1-800-540-3765

Package label

2	NDC:68788-4065-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	01/28/2026	
3	NDC:68788-4065-5	50 in 1 BOTTLE; Type 0: Not a Combination Product	01/28/2026	
4	NDC:68788-4065-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	01/28/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	01/28/2026	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-4065)

Revised: 1/2026

Preferred Pharmaceuticals Inc.