

EXTRA STRENGTH ACETAMINOPHEN- acetaminophen tablet
Preferred Pharmaceuticals Inc.

gc219

Active Ingredients

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 tablets (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.

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 tablets every 6 hours, as needed; not more than 6 tablets 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)

Inactive Ingredients

povidone, sodium starch glycolate, starch, stearic acid.

Questions or comments?

1-800-540-3765

Repackaged By: Preferred Pharmaceuticals Inc.

Extra Strength
Acetaminophen
500mg

Generic for Tylenol

Each tablet contains: Acetaminophen 500mg

Pkg Size: Exp Date: #####

Lot#: Batch#:


Ins:

Mfg: Geri-Care Pharm. Corp.

Prod#:


Warning

Liver Warning: This product contains acetaminophen. Severe liver damage may occur if you take more than 8 tablets (4,000mg of acetaminophen) in 24 hours. Do not use with any other drug containing acetaminophen (prescription or non prescription), 3 or more alcoholic drinks every day while using this product. Ask a doctor before use if you have liver disease or are taking the blood thinning drug warfarin. Keep out of reach of children. Store at 20° - 25°C (68° - 77°F). Tablet is round, white, imprinted with GC216



Directions English

Take tablet(s)
every hours.



GTIN

SN #####
EXP #####

Instrucciones Espanol:
Toma tableta(s)
cada horas.

Extra Strength Acetaminophen 500
mg
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Extra Strength Acetaminophen 500
mg
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Extra Strength Acetaminophen 500
mg
Qty: Ins:
Insurance NDC:
Lot: Bat:

Extra Strength Acetaminophen 500
mg
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Log

Chart

Billing

Patient

EXTRA STRENGTH ACETAMINOPHEN				
acetaminophen tablet				
Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source) NDC:68788-8873(NDC:57896-219)	
Route of Administration		ORAL		
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)			ACETAMINOPHEN	500 mg
Inactive Ingredients				
Ingredient Name				Strength
STARCH, CORN (UNII: O8232NY3SJ)				
CROSPVIDONE, UNSPECIFIED (UNII: 2S7830E561)				
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
Product Characteristics				
Color	white (WHITE)		Score	no score
Shape	ROUND (Round)		Size	12mm
Flavor			Imprint Code	GC;216
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-8873-2	20 in 1 BOTTLE; Type 0: Not a Combination Product	05/16/2025	

2	NDC:68788-8873-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	05/16/2025	
3	NDC:68788-8873-5	50 in 1 BOTTLE; Type 0: Not a Combination Product	05/16/2025	
4	NDC:68788-8873-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/16/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	05/16/2025	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-8873)