ACETAMINOPHEN- acetaminophen tablet Preferred Pharmaceuticals Inc.

5427-Major

Drug Facts

Active Ingredient (in each tablet)

Acetaminophen 500 mg

Purpose

Pain reliever/ Fever reducer

Uses

Temporarily reduces fever and relieves minor aches and pains due to:

- headache □
- muscular aches □
- backache □
- minor pain of arthritis □
- common cold □
- toothache □
- premenstrual and
- menstrual cramps

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take

- more than 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.

- 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 \square

- 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.

Overdose warning: In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222) 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 (see overdose warning)

adults and children 12 years and over:

- take 2 tablets every 6 hours while symptoms last
- do not take more than 6 tablets in 24 hours, unless directed by a doctor
- do not take for more than 10 days unless directed by a doctor

children under 12 years: ask a doctor

Other Information

store at room temperature

Povidone, pregelatinized starch, sodium starch glycolate, stearic acid

Questions or comments?

Call (800) 616-2471

SAVE CARTON FOR COMPLETE DRUG FACTS

Do not use if carton is open or if imprinted safety seal under cap is broken or missing.

Distributed By:

MAJOR® PHARMACEUTICALS

17177 N Laurel Park Drive, Suite 233

Livonia, MI 48152

*This product is not manufactured or distributed by McNeil Consumer Healthcare, owner of the registered trademark Tylenol®

MAJOR

THIS LARGE SIZE PACKAGE FOR DISPENSING PURPOSE ONLY

See New Warnings Information and Directions

Acetaminophen Extra Strength Tablets

Aspirin Free

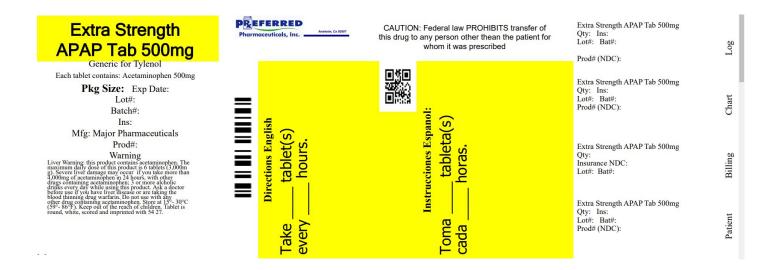
Extra Strenght Pain reliever

Compare to active ingredient in Extra Strength Tylenol®* Tablets

500mg each

Relabeled By: Preferred Pharmaceuticals Inc.

NDC 68788-8636



ACETAMINOPHEN

acetaminophen tablet

Pı	roduct Information	ct Information				
Pr	oduct Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68788-8636(NDC:0904-6730)		
Ro	oute of Administration	ORAL				

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	500 mg			

Inactive Ingredients				
Ingredient Name	Strength			
STARCH, CORN (UNII: O8232NY3SJ)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
POVIDONE K30 (UNII: U725QWY32X)				

SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)

Product Characteristics					
Color	white	Score	2 pieces		
Shape	ROUND	Size	12mm		
Flavor		Imprint Code	54;27		
Contains					

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788- 8636-2	20 in 1 BOTTLE; Type 0: Not a Combination Product	04/17/2024	
2	NDC:68788- 8636-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	04/17/2024	
3	NDC:68788- 8636-5	50 in 1 BOTTLE; Type 0: Not a Combination Product	04/17/2024	
4	NDC:68788- 8636-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	04/17/2024	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	343	04/17/2024		

Labeler - Preferrred Pharmaceuticals Inc. (791119022)

Registrant - Preferrred Pharmaceuticals Inc. (791119022)

Establishment					
Name	Address	ID/FEI	Business Operations		
Preferrred Pharmaceuticals Inc.		791119022	REPACK(68788-8636)		

Revised: 4/2024 Preferrred Pharmaceuticals Inc.