#### ACETAMINOPHEN- acetaminophen tablet Preferred Pharmaceuticals Inc.

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#### **Major Pharmaceuticals Acetaminophen Drug Facts**

#### Active ingredient (in each caplet)

Acetaminophen 500 mg

#### Purpose

Pain reliever/fever reducer

#### Uses

- temporarily relieves minor aches and pains due to:
- the common cold
- headache
- backache
- minor pain of arthritis
- toothache
- muscular aches
- premenstrual and menstrual cramps
- temporarily reduces fever

#### Warnings

**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.

#### 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 have ever had an allergic reaction to this product or any of its ingredients

# 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 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	<ul> <li>take 2 caplets every 6 hours while symptoms last</li> <li>do not take more than 6 caplets in 24 hours, unless directed by a doctor</li> <li>do not use for more than 10 days unless directed by a doctor</li> </ul>				
children under 12 years ask a doctor					

# Other information

• store at 20-25°C (68-77°F)

# Inactive ingredients

carnauba wax, corn starch\*, croscarmellose sodium\*, hypromellose, polyethylene

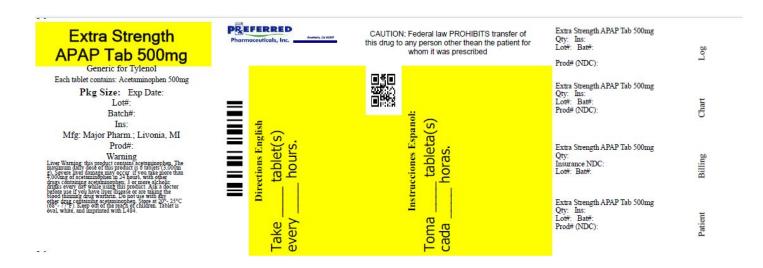
glycol, povidone, pregelatinized starch, sodium starch glycolate\*, stearic acid \*may contain one or more of these ingredients

Questions or comments? 1-800-616-2471

#### **Principal Display Panel**

EXTRA STRENGTH Acetaminophen PAIN RELIEVER/FEVER REDUCER ASPIRIN FREE FOR ADULTS CAPLETS Compare to the active ingredient in Extra Strength Tylenol® Caplets 500 mg. Each

### **Repackaged By: Preferred Pharmaceuticals Inc.**



ACETAMINOPHEN acetaminophen tablet						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68788-8070(NDC:0904-6720)			
Route of Administration	ORAL					
Active Ingredient/Active Moiety						

		lr	ngredient Nam	e		<b>Basis of</b>	Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNI				NOPHEN - UNII:3620	9ITL9D)	ACETAMINO	PHEN	500 mg	
In	nactive Ingre	dients							
Ingredient Name								trength	
	TARCH, CORN (U								
			ED (UNII: 3NXW29)						
				3WJQ0SDW1A)					
	DVIDONE, UNSPE								
	FEARIC ACID (UN		•	0)					
Cr	KUSCARMELLUS		(UNII: M28OL1HH4	0)					
P	roduct Chara	acteristi	rs						
_	olor		WHITE	Score			no score		
	hape		OVAL	Size			16mm		
	avor			Imprint Code			L484		
Са	ontains								
Co	ontains								
Co	ontains								
	ontains ackaging								
Pa	ackaging		Package Desc	ription		ing Start ate		ting End ate	
Pa #	ackaging		Package Desc DTTLE; Type 0: Not	•		-			
Pa	ackaging Item Code NDC:68788-	20 in 1 BO Product	-	a Combination	D	-			
<b>P</b> a # 1	ackaging Item Code NDC:68788- 8070-2 NDC:68788-	20 in 1 BO Product 30 in 1 BO Product 50 in 1 BO Product	OTTLE; Type 0: Not OTTLE; Type 0: Not OTTLE; Type 0: Not	a Combination a Combination a Combination	D 08/23/2021	-			
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<b>P</b> # 1 2 3	ackaging Item Code NDC:68788- 8070-2 NDC:68788- 8070-3 NDC:68788- 8070-5 NDC:68788-	20 in 1 BO Product 30 in 1 BO Product 50 in 1 BO Product 100 in 1 B	OTTLE; Type 0: Not OTTLE; Type 0: Not OTTLE; Type 0: Not	a Combination a Combination a Combination	D. 08/23/2021 08/23/2021 08/23/2021	-			
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Labeler - Preferred Pharmaceuticals Inc. (791119022)

**Registrant -** Preferred Pharmaceuticals Inc. (791119022)

Establishment							
Name	Address	ID/FEI	<b>Business Operations</b>				
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-8070)				