

EXTRA STRENGTH PAIN RELIEVER- acetaminophen tablet
INGLES MARKETS, INC.

LL222

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. The maximum daily dose of this product is 6

caplets in 24 hours. 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.

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 of age and over	take 1-2 caplets every 4-6 hours, as needed. Do not take more than 6 caplets in 24 hours.
Children under 12 years	Ask a doctor

Other Information

- **TAMPER EVIDENT: Do not use if imprinted seal under cap is missing or broken.**
- store at 20°C-25°C (68°F-77°F)

Inactive ingredients

povidone, sodium starch glycolate, starch, stearic acid

Package label



EXTRA STRENGTH Pain Reliever

ACETAMINOPHEN 500 mg

NON-ASPIRIN

- pain reliever
- fever reducer



100 CAPLETS

Compare to Active Ingredient in Extra Strength Tylenol®*



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NON-ASPIRIN
EXTRA STRENGTH
Pain Reliever
 ACETAMINOPHEN 500 mg



Drug Facts

SAVE CARTON FOR COMPLETE DRUG FACTS

Active ingredient (in each caplet)

Acetaminophen 500 mg, Pain Reliever / Fever Reducer

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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.

Drug Facts (continued)

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.

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Children under 12 years	Ask a doctor.

Other information • TAMPER EVIDENT: Do not use if imprinted seal under cap is missing or broken.

- store at 20°C - 25°C (68°F-77°F)

Inactive ingredients: povidone, sodium starch glycolate, starch, stearic acid.

* This product is not manufactured or distributed by the owner of the registered trademark TYLENOL®.

REV 221-0919

DISTRIBUTED BY: **ingles**
P.O. Box 6676, ASHEVILLE, NC 28816



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EXTRA STRENGTH PAIN RELIEVER

acetaminophen tablet

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:62936-0222

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		
	POVIDONE (UNII: FZ989GH94E)			
	SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)			
	STARCH, CORN (UNII: O8232NY3SJ)			
	STEARIC ACID (UNII: 4ELV7Z65AP)			
Product Characteristics				
Color	white (WHITE)	Score	no score	
Shape	CAPSULE (Capsule)	Size	17mm	
Flavor		Imprint Code	AZ 328	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62936-0222-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2019	
2	NDC:62936-0222-5	50 in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2019	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M013	08/01/2018		

Labeler - INGLES MARKETS, INC. (024418584)

Registrant - Geri-Care Pharmaceutical Corp (611196254)