ACETAMINOPHEN- acetaminophen tablet
NuCare Pharmaceuticals, Inc.

Acetaminophen Tablets 500 mg

ACTIVE INGREDIENT (IN EACH CAPLET)

Acetaminophen, USP 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 24 hours.
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks everyday 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 drugs 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 other 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 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

Over dose warning: In case of overdose, get medical help or Contact 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 caplets every 6 hours while symptoms last • do not take more than 6 caplets in 24 hours, unless directed by a doctor • do not use for more than 10 days unless directed by a doctor
children under 12 years	• ask a doctor

OTHER INFORMATION

•store at 20-25°C (68-77°F). See USP Controlled Room Temparature

•avoid high humidity

•See end panel for lot number and expiration date

INACTIVE INGREDIENTS

hydroxyethyl methyl cellulose, polyethylene glycol, povidone, pregelatinized starch, sodium starch glycolate, stearic acid.

QUESTIONS OR COMMENTS ?

Contact **1-877-770-3183**Mon-Fri 8:00 AM EST to 5:00 PM PST.

+All trademarkes are property of their reapective owners.

This product is not affliated with the makers/owners of

Extra Strength Tylenol ®Caplets.

Distributed by:

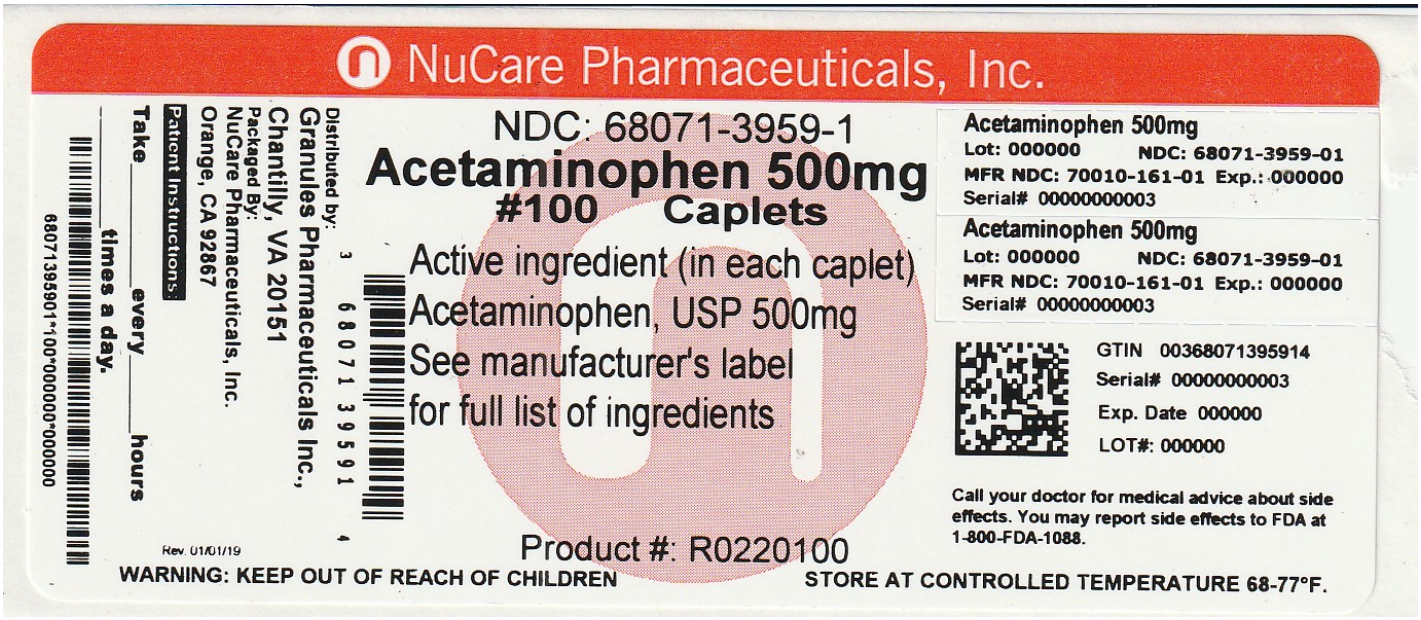
Granules Pharmaceuticals INc.,

Chantilly, VA 20151

MADE IN INDIA

Rev.08/21

PRINCIPAL DISPLAY PANEL



ACETAMINOPHEN

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68071-3959(NDC:70010-161)
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 K30 (UNII: U725QWY32X)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	

Product Characteristics			
Color	white (White to off-White)	**Score**	no score
Shape	CAPSULE (Caplet shape)	**Size**	17mm
Flavor		**Imprint Code**	G551
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-3959-1	1 in 1 CARTON	01/27/2026	
1		100 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	02/15/2022	

Labeler - NuCare Pharmaceuticals, Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals, Inc.		010632300	relabel(68071-3959)

Revised: 1/2026 NuCare Pharmaceuticals, Inc.