ACETAMINOPHEN- acetaminophen tablet
Trifecta Pharmaceuticals USA LLC

Acetaminophen 500mg

Active Ingredient

Acetaminophen USP, 500mg (in each tablet)

Purpose

Pain Releiver / Fever Reducer

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 are allergic to acetaminophen or any of the inactive ingredients in this product.

Uses

Temporarily relieves minor aches and pains due to

Common cold

headache

minor pain of arthritis

toothache

muscular aches

premenstrual and menstrual cramps

temporarily relieves fever

Warnings

Liver warning

This product contains acetaminophen. Sever liver damage may occur if you take

- more than 4000mg 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 ad seek medical medical help right away

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.

Keep out of reach of children

Keep out of reach of children

Directions

Do not take more than directed. See overdose warning

Adults and children 12 years and older
- 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 use for more than 10 days unless directed by a doctor

Children under 12 years of age
- Ask a doctor

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 a sign of a serious condition

If pregnant or breast-feeding

Ask a health professional before use

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 cretical for adults as well as children even if you do not notice symptoms.

Other Information

- Store at 20ºC - 25ºC (68ºF - 77ºF)
- Avoid high humidity
- See end panel for expiration date and lot number

Inactive Ingredients

hydroxypropyl methyl cellulose, polyethylene glycol, povidone K-30, Pregelatinized starch, Sodium starch glycoate, Stearic Acid

Questions or Comments?

Call 1-888-296-9067

Distributed By:

Distributed By:

Trifecta Pharmaceuticals USA®

Ft. Lauderdale, FL. 33301 USA

1-888-296-9067

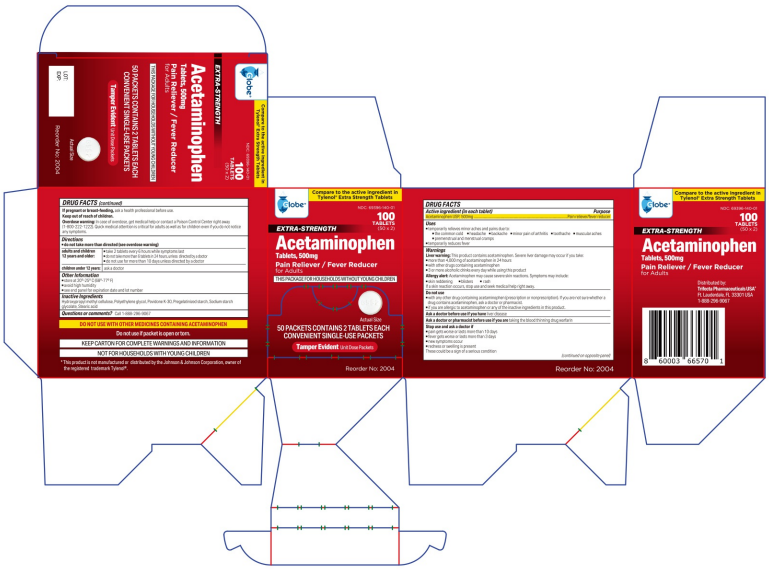
*This product is not manufactured or distributed by the Johnson & Johnson Corporation, owner of the registered trademark Tylenol®

Reorder No. 2004

Made in the USA

Label

OUTSIDE BOX



ACETAMINOPHEN

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69396-140
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**
STEARIC ACID (UNII: 4ELV7Z65AP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	

STARCH, CORN (UNII: O8232NY3SJ)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POVIDONE K30 (UNII: U725QWY32X)	

Product Characteristics

Color	white	**Score**	no score
Shape	ROUND	**Size**	12mm
Flavor		**Imprint Code**	G552
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69396-140-03	10000 in 1 BAG	08/08/2023	
1		2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:69396-140-01	50 in 1 BOX	08/08/2023	
2		2 in 1 PACKET; 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	08/08/2023	

Labeler - Trifecta Pharmaceuticals USA LLC (079424163)

Revised: 6/2024 Trifecta Pharmaceuticals USA LLC