ACETAMINOPHEN EXTRA STRENGTH- acetaminophen tablet, coated United Natural Foods, Inc. dba UNFI

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

1004-ELN-2022-0915

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 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

- 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 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 between 20-25°C (68-77°F)
- retain carton for complete product information

Inactive ingredients

hypromellose, mineral oil, polyethylene glycol, polysorbate 80, povidone, pregelatinized starch, sodium starch glycolate, stearic acid, titanium dioxide

Questions or comments?

1-855-423-2630

PRINCIPAL DISPLAY PANEL

Equaline®

NDC 41163-500-01

compare to Extra Strength Tylenol® Caplets active ingredient*

extra strength

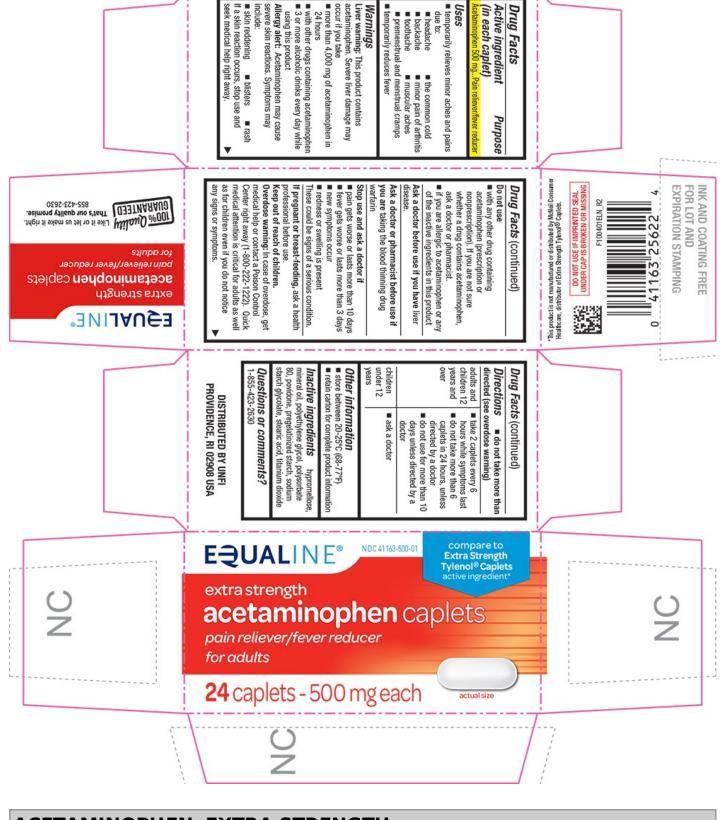
acetaminophen caplets

pain reliever/fever reducer

for adults

24 caplets - 500 mg each

actual size



ACETAMINOPHEN EXTRA STRENGTH

acetaminophen tablet, coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41163-500
Route 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			
HYPROMELLOSES (UNII: 3NXW29V3WO)				
MINERAL OIL (UNII: T5L8T28FGP)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
POLYSORBATE 80 (UNII: 6OZP39ZG8H)				
POVIDONE (UNII: FZ989GH94E)				
STARCH, PREGELATINIZED CORN (UNII: O8232NY3SJ)				
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				

Product Characteristics			
Color	white	Score	no score
Shape	OVAL	Size	17mm
Flavor		Imprint Code	M2A4;57344
Contains			

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:41163- 500-01	1 in 1 CARTON	11/01/2013		
1		24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			
2	NDC:41163- 500-02	1 in 1 CARTON	11/01/2013	05/31/2020	
2		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			
3	NDC:41163- 500-04	1 in 1 CARTON	11/01/2013		
3		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			
4	NDC:41163- 500-05	1 in 1 CARTON	11/01/2013	12/01/2019	
4		250 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			
5	NDC:41163- 500-06	1 in 1 CARTON	11/01/2013		
5		500 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			
6	NDC:41163- 500-17	1 in 1 CARTON	11/01/2013	03/31/2018	
6		150 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part343	11/01/2013			

Labeler - United Natural Foods, Inc. dba UNFI (943556183)

Revised: 5/2023 United Natural Foods, Inc. dba UNFI