FIRST AID ONLY EXTRA-STRENGTH NON-ASPIRIN- acetaminophen tablet, film coated Acme United Corporation

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.

First Aid Only Extra-Strength Non-Aspirin

Drug Facts

Active ingredient (in each tablet)

Acetaminophen 500 mg

Purpose

Pain reliever/fever reducer

Uses

For the temporary relief of minor aches and pains associated with

- headache
- toothache
- minor arthritis pain
- muscular aches
- common cold
- menstrual cramps

temporarily reduces fever

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 8 tablets in 24 hours, which is the maximum daily amount
- 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.
- for more than 10 days for pain unless directed by a doctor
- for more than 3 days for fever unless directed by a doctor

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 using and ask a doctor if

- symptoms do not improve
- pain or fever persists or gets worse
- new symptoms occur
- redness or swelling is present

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

In case of accidental overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

• do not use more than directed

Adults and children: (12 years and older)

Take 2 tablets every 4 to 6 hours as needed. Do not take more than 8 tablets in 24 hours.

Children under 12 years:

Do not give this adult strength product to children under 12 years of age; this will provide more than the recommended dose (overdose) and may cause liver damage.

Other information

- store at room temperature 59°-86°F (15°-30°C)
- tamper-evident sealed packets
- do not use any opened or torn packets

Inactive ingredients

corn starch, hypromellose, maltodextrin*, microcrystalline cellulose*, polyethylene glycol, povidone*, pregelatinized corn starch*, sodium starch glycolate*, stearic acid, titanium dioxide*

* May contain

Questions? 1-800-835-2263

First Aid Only Extra-Strength Non-Aspirin Label

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Manufactured for:
Acme United Corporation
www.FirstAidOnly.com
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Covered by one or more of US Patent Numbers:
D495,951 S; D495,952 S; D495,953 S
B0X7008-revC



FIRST AID ONLY EXTRA-STRENGTH NON-ASPIRIN

acetaminophen tablet, film coated

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0924-0228(NDC:47682-125)	
Route of Administration	ORAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D)	ACETAMINOPHEN	500 mg		

Inactive Ingredients				
Ingredient Name	Strength			
CELLULOSE, MICRO CRYSTALLINE (UNII: OP1R32D61U)				
HYPROMELLOSES (UNII: 3NXW29V3WO)				
MALTO DEXTRIN (UNII: 7CVR7L4A2D)				
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)				
PO VIDO NE (UNII: FZ989 GH94E)				
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	ROUND (ROUND)	Size	12mm
Flavor		Imprint Code	FR;33
Contains			

	Packaging			
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:0924-0228-01	20 in 1 BOX, UNIT-DOSE	12/30/2008	
	1	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 not final	part343	12/30/2008		

Labeler - Acme United Corporation (001180207)

Registrant - Acme United Corporation (001180207)

Establishment			
Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	relabel(0924-0228), repack(0924-0228)

Establishment				
Name	Address	ID/FEI	Business Operations	
Acme United Corporation		080119599	relabel(0924-0228), repack(0924-0228)	

Revised: 3/2018 Acme United Corporation