ARAMARK EXTRA STRENGTH ACETAMINOPHEN- acetaminophen tablet
Western First Aid Safety DBA Aramark

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.

Aramark Extra Strength Acetaminophen

Drug Facts

Active Ingredient (in each tablet)
Acetaminophen 500 mg

Purpose
Acetaminophen .....Pain Reliever/Fever Reducer

Uses:
Temporarily Relieves minor aches and pains due to:
• Headache, toothaches, menstrual cramps, the common cold,
• muscular aches, minor pain of arthritis
Temporarily reduces fever

Warnings:
Allergy Alert: Acetaminophen may cause a severe skin reactions.
Symptoms may include:
• Skin reddening •Blisters •Rash
• If a skin reaction occurs, stop use and seek medical right away.

Liver Warning: This product contains an acetaminophen. Severe liver damage may occur if you take:
• more than 8 tablets in 24 hours, which is the maximum daily amount
• 3 or more alcoholic drinks every day while using this product

• take more or for a longer time than directed

Do not use:
• if you have ever had an allergic reaction to any other pain
• reliever/fever reducer
• right before or after heart surgery
• if you have ever had an allergic reaction to this product or any of its ingredients

Ask a doctor before use if:
• you have liver disease

Ask a doctor or pharmacist before use if you are:
• under a doctor's care for any serious condition
• taking the blood thinning drug Warfarin

When using this product:
• take with food or milk if stomach upset occurs
• the risk of heart attack or stroke may increase if you use more than directed or for longer than directed

Stop use and ask a doctor if:
• pain gets worse lasts more than 10 days
• feel faint
• pain gets worse or lasts more than 10 days
• fever gets worse or lasts more than 3 days
• redness or swelling is present in the painful area
• any new symptoms occur

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. 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 take more than directed**
• adults and children 12 years and over: take 2 tablets every 4 to 6 hours as needed
• do not exceed 8 tablets in 24 hours, unless directed by a doctor
• Children under 12 years: **Do not use this product**

Other information:
read all warnings and directions before use. Keep carton. Store at 20-25°C (68-77°F) avoid excessive heat above 40°C (104°F)

Inactive Ingredients:
Microcrystalline Cellulose, Povidone, Sodium Starch Glycolate, Starch, Stearic Acid

Package Labeling

aramark

100 TABLETS PER BOX **Part # 90433B**

Extra Strength
Acetaminophen
Pain Reliever/Fever Reducer

Temporary relief of minor aches and pains associated with a cold, headache, tootache, muscular aches, and for the reduction of fever

Compare active ingredient to:
Extra Strength Tylenol®
Registered Trademark of Johnson & Johnson

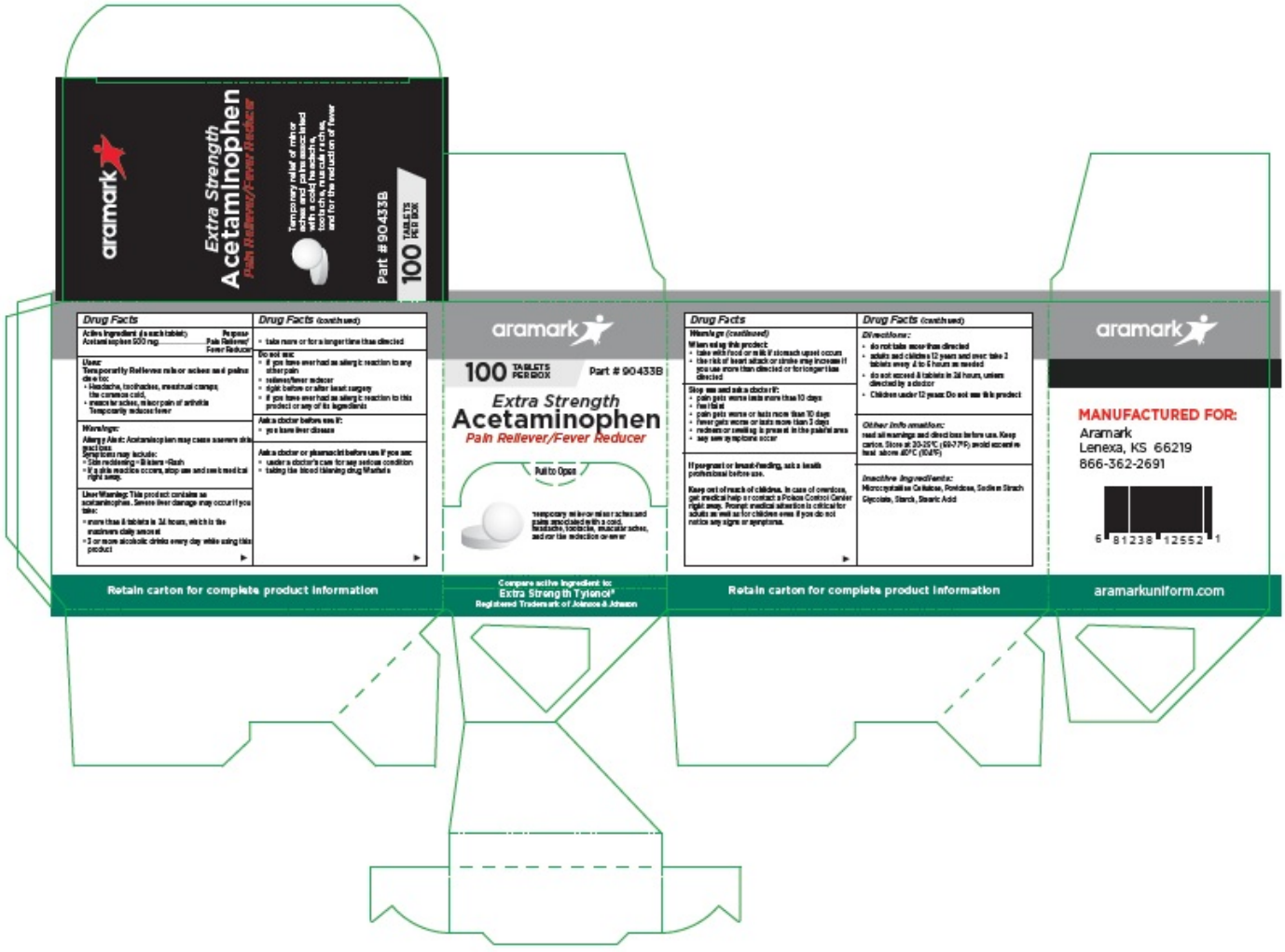
MANUFACTURED FOR:

Aramark
Lenexa, KS 66219
866-362-2691

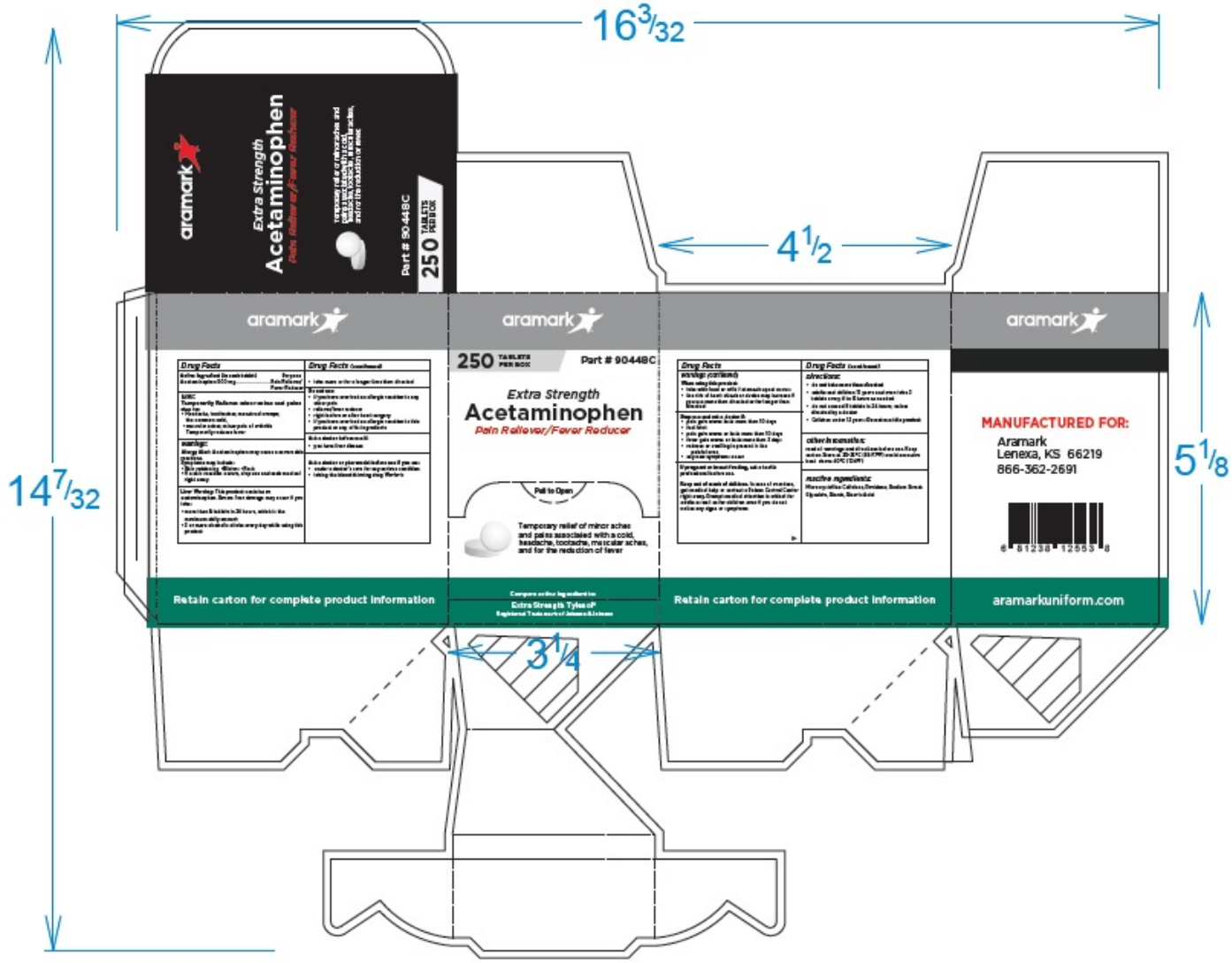
aramarkuniform.com

Retain carton for complete product information

100 Tablet Box



250 Tablet Box



2-Tablet Packet

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ARAMARK EXTRA STRENGTH ACETAMINOPHEN

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81238-3500
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**
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POVIDONE (UNII: FZ989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	white	**Score**	no score
Shape	ROUND	**Size**	18mm
Flavor		**Imprint Code**	FR1
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81238-3500-1	50 in 1 BOX	05/14/2021	
1		2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:81238-3500-2	125 in 1 BOX	05/14/2021	
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 not final	part343	05/14/2021	

Labeler - Western First Aid Safety DBA Aramark (043861524)

Registrant - Western First Aid Safety DBA Aramark (043861524)

Establishment

Name	Address	ID/FEI	Business Operations
ULTRA SEAL CORPORATION		085752004	pack(81238-3500)

Establishment

Name	Address	ID/FEI	Business Operations
ULTRA SEAL CORPORATION		944090448	manufacture(81238-3500)

Revised: 7/2021 Western First Aid Safety DBA Aramark