MEDI-FIRST PLUS EXTRA STRENGTH NON-ASPIRIN- acetaminophen tablet, film coated

MEDI-FIRST EXTRA STRENGTH NON-ASPIRIN- acetaminophen tablet, film coated

MEDIQUE EXTRA STRENGTH APAP- acetaminophen tablet, film coated Unifirst First Aid Corporation

Acetaminophen 500 mg

Drug Facts

Active Ingredient (in each tablet)

Acetaminophen 500 mg

Purpose

Pain reliever / fever reducer

Uses

temporarily relieves minor aches and pains due to:

- headache
- muscular aches
- backache
- minor pain of arthritis
- the common cold
- toothache
- 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 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.
- 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 using and ask a doctor if

- pain gets worse or lasts more than for 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.

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 use more than directed (see overdose warning)

Adults and children 12 years and over

- 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

Ask a doctor

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*, polyethylene glycol*, povidone (K-30)*, pregelatinized starch*, purified water*, sodium starch glycolate, stearic acid, titanium dioxide*

* May contain

Medique

Extra Strength APAP

Non-Aspirin Tablets

THIS PACKAGE IS FOR HOUSEHOLDS WITHOUT YOUNG CHILDREN.

Pain Reliever / Fever Reducer • Acetaminophen 500 mg

Compare active ingredient to Tylenol®

Registered McNeil Products

Tamper Evident Unit Dose Packets

24 Tablets

 (12×2)



MEDI-FIRST PLUS EXTRA STRENGTH NON-ASPIRIN

acetaminophen tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47682-274
Route of Administration	ORAL		

Active Ingredient/Active Moiety

ı	Ingredient Name	Basis of Strength	Strength
ı	ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII: 36209ITL9D)	ACETAMINOPHEN	500 mg

Inactive Ingredients		
Ingredient Name	Strength	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)		
STARCH, CORN (UNII: O8232NY3SJ)		
POVIDONE K30 (UNII: U725QWY32X)		

Product Characteristics				
Color white ((White to Off-White)) Score no score				
Shape	ROUND	Size	12mm	
Flavor		Imprint Code	G552	
Contains				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:47682-274- 33	50 in 1 BOX	03/28/2024		
1		2 in 1 PACKET; Type 0: Not a Combination Product			
2	NDC:47682-274- 48	125 in 1 BOX	03/28/2024		
2		2 in 1 PACKET; Type 0: Not a Combination Product			

Marketing Information			
Marketing Application Number or Monograph Marketing Start Marketing End Category Citation Date Date			
OTC Monograph Drug	M013	03/28/2024	

MEDI-FIRST EXTRA STRENGTH NON-ASPIRIN

acetaminophen tablet, film coated

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:47682-273 Route of Administration ORAL

Active Ingredient/Active Moiety		
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ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII: 36209ITL9D)	ACETAMINOPHEN	500 mg

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POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
POVIDONE K30 (UNII: U725QWY32X)			
STEARIC ACID (UNII: 4ELV7Z65AP)			

Product Characteristics				
Color	white ((White to Off-White))	Score	no score	
Shape	ROUND	Size	12mm	
Flavor		Imprint Code	G552	
Contains				

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:47682-273- 13	250 in 1 BOX	03/28/2024			
1	NDC:47682-273- 99	2 in 1 PACKET; Type 0: Not a Combination Product				
2	NDC:47682-273- 33	50 in 1 BOX	03/28/2024			
2	NDC:47682-273- 99	2 in 1 PACKET; Type 0: Not a Combination Product				
3	NDC:47682-273- 48	125 in 1 BOX	03/28/2024			
3	NDC:47682-273- 99	2 in 1 PACKET; Type 0: Not a Combination Product				
4	NDC:47682-273- 50	25 in 1 BOX	03/28/2024			
4	NDC:47682-273- 99	2 in 1 PACKET; Type 0: Not a Combination Product				

Marketing In	formation		
Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC Monograph Drug	M013	03/28/2024	

MEDIQUE EXTRA STRENGTH APAP

acetaminophen tablet, film coated

Produ	ct In	form	ation
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:47682-272

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name

ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)

ACETAMINOPHEN

ACETAMINOPHEN

ACETAMINOPHEN

500 mg

Inactive Ingredients

Ingredient Name
Strength

STEARIC ACID (UNII: 4ELV7Z65AP)

HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)

SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)

POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)

STARCH, CORN (UNII: 08232NY3SJ)

POVIDONE K30 (UNII: U725QWY32X)

Product Characteristics			
Color	white ((White to Off-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:47682-272- 13	250 in 1 BOX	03/28/2024	
1	NDC:47682-272- 99	2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:47682-272- 33	50 in 1 BOX	03/28/2024	
2	NDC:47682-272- 99	2 in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:47682-272- 48	125 in 1 BOX	03/28/2024	
3	NDC:47682-272- 99	2 in 1 PACKET; Type 0: Not a Combination Product		
	NDC: 47602 272			

4 64	4	12 in 1 BOX	03/28/2024	
4 NI		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
отс	Monograph Drug	M013	03/28/2024	

Labeler - Unifirst First Aid Corporation (832947092)

Establishment			
Name	Address	ID/FEI	Business Operations
Prestige Packaging		080667761	pack(47682-272, 47682-273, 47682-274)

Establishment				
Name	Address	ID/FEI	Business Operations	
Medique Products		086911794	pack(47682-272, 47682-273, 47682-274)	

Revised: 3/2024 Unifirst First Aid Corporation