

MEDIQUE APAP EXTRA STRENGTH- acetaminophen tablet, film coated
MEDI-FIRST NON-ASPIRIN EXTRA STRENGTH- acetaminophen tablet, film coated
MEDI FIRST PLUS NON-ASPIRIN EXTRA STRENGTH- 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
- minor pain of arthritis
- backache
- 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 use and ask a doctor if

- pain gets worse or lasts for more than 10 days
- fever gets worse or lasts for 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 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*

Questions or comments? 1-800-634-7680

Medi-First Extra Strength Non Aspirin

Medi-First®

Extra Strength

Non-Aspirin

Pain Reliever/Fever Reducer

Aches, Fever ● Acetaminophen 500mg

Pull to Open

Compare active ingredient to:

Extra Strength Tylenol®

Registered Trademark of McNeil Consumer Products

This Package is for Households without Young Children.

Tamper Evident Unit Dose Packets

100 Tablets

(50 x 2)



Medi First Plus Extra Strength Non-Aspirin

Medi First®

Plus

Extra Strength

Non-Aspirin

Acetaminophen 500 mg

Pull To Open

This Package is For Households Without Young Children.

Pain Reliever/Fever Reducer

Compare Active ingredient to:

Extra Strength Tylenol®

Registered Trademark of McNeil Consumer

Tamper Evident Unit Dose Packets

100 Tablets

(50 x 2's)

Drug Facts
Active ingredient (in each tablet) **Purpose**
 Acetaminophen 500 mg Pain reliever/fever reducer

Uses
 Temporarily relieves minor aches and pains due to:
 ■ headache ■ muscular aches ■ minor pain of arthritis ■ backache
 ■ 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

Always alert: Acetaminophen may cause severe skin reactions. Symptoms may include:
 ■ skin redness ■ 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 (continued on opposite panel)

Retain carton for complete product information

MEDI FIRST Plus Extra Strength Non-Aspirin
 Acetaminophen 500 mg
 Pull To Open
 THIS PACKAGE IS FOR HOUSEHOLDS WITHOUT YOUNG CHILDREN
 Pain Reliever/Fever Reducer
 Compare active ingredient to:
 Extra Strength Tylenol®
 Registered Trademark of McNeil Consumer
 Tamper Evident Unit Dose Packets

Drug Facts (continued)
 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 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.

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 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 8 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

Drug Facts (continued)
Inactive ingredients
 corn starch, hydroxybenzoin, polyethylene glycol, povidone (K-30), pregelatinized starch, purified water, sodium starch glycolate, stearic acid, titanium dioxide

*may contain

Questions or comments? 1-800-634-7680

Barcode: 4768217564

Border #99433

Manufactured for Medique Products
 Fort Myers, FL 33967 USA
 1-800-634-7680
 www.medicueproducts.com

Medique Extra Strength APAP

Medique®

Extra Strength

APAP

Non-Aspirin Tablets

This Package is For Households Without Young Children

Pain Reliever/Fever Reducer ● Acetaminophen 500mg

Compare active ingredient to Tylenol®

Registered McNeil Consumer Products

Tamper Evident Unit Dose Packets

Drug Facts (continued)
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 8 tablets in 24 hours, unless directed by a doctor
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 ■ tamper-evident sealed packets
 ■ do not use any opened or torn packets

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Drug Facts
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 ■ 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

Always alert: Acetaminophen may cause severe skin reactions. Symptoms may include:
 ■ skin redness ■ blisters ■ rash
 If a skin reaction occurs, stop use and seek medical help right away.

(continued on inside panel)

Retain carton for complete product information

Medique® Extra Strength APAP Non-Aspirin Tablets
 THIS PACKAGE IS FOR HOUSEHOLDS WITHOUT YOUNG CHILDREN
 Pain Reliever/Fever Reducer • Acetaminophen 500 mg
 24 Tablets (12 x 2)
 Compare active ingredient to Tylenol®
 Registered Trademark of McNeil Consumer Products
 Tamper Evident Unit Dose Packets

Barcode: 4768217564

Border #17564

Manufactured for Medique Products
 Fort Myers, FL 33967
 1-800-634-7680
 www.medicueproducts.com

Drug Facts (continued)
 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
 ■ if you are taking the blood thinning drug warfarin

Stop use and ask a doctor if:
 ■ pain gets worse or lasts 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.

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.

MEDIQUE APAP EXTRA STRENGTH

acetaminophen tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47682-175
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg
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Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STARCH, CORN (UNII: O8232NY3SJ)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white (white)	Score	no score
Shape	ROUND (ROUND)	Size	12mm
Flavor		Imprint Code	AZ;235
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-175-99	2 in 1 PACKET; Type 0: Not a Combination Product	12/30/2008	
2	NDC:47682-175-13	250 in 1 BOX	12/30/2008	
2		2 in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:47682-175-33	50 in 1 BOX	12/30/2008	
3		2 in 1 PACKET; Type 0: Not a Combination Product		
4	NDC:47682-175-48	125 in 1 BOX	12/30/2008	
4		2 in 1 PACKET; Type 0: Not a Combination Product		
5	NDC:47682-175-64	12 in 1 BOX	12/30/2008	
5		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	12/30/2008	

MEDI-FIRST NON-ASPIRIN EXTRA STRENGTH

acetaminophen tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47682-804
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
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white (white)	Score	no score
Shape	ROUND (ROUND)	Size	12mm
Flavor		Imprint Code	AZ;235
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-804-33	50 in 1 BOX	12/30/2008	
1		2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:47682-804-48	125 in 1 BOX	12/30/2008	
2		2 in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:47682-804-13	250 in 1 BOX	12/30/2008	
3		2 in 1 PACKET; Type 0: Not a Combination Product		
4	NDC:47682-804-99	2 in 1 PACKET; Type 0: Not a Combination Product	12/30/2008	
5	NDC:47682-804-50	25 in 1 BOX	12/30/2008	
5		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	12/30/2008	

MEDI FIRST PLUS NON-ASPIRIN EXTRA STRENGTH

acetaminophen tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47682-904
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
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white (white)	Score	no score
Shape	ROUND (ROUND)	Size	12mm
Flavor		Imprint Code	AZ;235
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-904-33	50 in 1 BOX	12/30/2008	
1		2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:47682-904-48	125 in 1 BOX	12/30/2008	

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	12/30/2008	

Labeler - Unifirst First Aid Corporation (832947092)

Revised: 9/2025

Unifirst First Aid Corporation