

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 x 2)



MEDI-FIRST®

PLUS

100 TABLETS

(50X 2'S)

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

MEDI-FIRST®
Extra Strength
Non-Aspirin
100
Tablets
(50x2)
Pain Reliever/Fever Reducer
✓Aches, Fever • Acetaminophen 500 mg
Pull to Open
Compare active ingredient to:
Extra Strength Tylenol®
Registered Trademark of McNeil Consumer Products
THIS PACKAGE IS FOR HOUSEHOLDS
WITHOUT YOUNG CHILDREN

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

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

Inactive Ingredients

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

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:47682-274-33	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 Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End 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
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Route of Administration ORAL

Active Ingredient/Active Moiety

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Inactive Ingredients

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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:47682-273-13	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		
5	NDC:47682-273-99	2 in 1 PACKET; Type 0: Not a Combination Product	03/28/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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MEDIQUE EXTRA STRENGTH APAP

acetaminophen tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47682-272
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
STEARIC ACID (UNII: 4ELV7Z65AP)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
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			

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		
4	NDC:47682-272-64	12 in 1 BOX	03/28/2024	

4	NDC:47682-272-99	2 in 1 PACKET; Type 0: Not a Combination Product		
5	NDC:47682-272-99	2 in 1 PACKET; Type 0: Not a Combination Product	03/28/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC 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: 9/2025

Unifirst First Aid Corporation