

MEDIQUE APAP ACETAMINOPHEN- medique apap acetaminophen tablet, coated

HF Acquisition Co LLC, DBA HealthFirst

Drug Facts

Active ingredient (in each tablet)

Acetaminophen 325 mg

Pain reliever/fever reducer

Uses

temporarily relieves minor aches and pains due to:

headache

muscular aches

minor arthritis pain

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 using and ask a doctor if

pain gets worse and 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.

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

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Directions

do not use more than directed (see overdose warning)

Adults and children: (12 years and over)

take 2 tablets every 4 to 6 hours while symptoms last.

do not take more than 10 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, sodium starch glycolate, stearic acid, titanium dioxide*

May contain*

Pain Reliever/Fever Reducer • Acetaminophen 325 mg in a pouch NDC (51662-1273-2)

(01)00351662127322
(17)230919
(21)351662141212
(10)HF1234TESTING

SEE MANUFACTURE'S INSERT
DISTRIBUTED BY HF ACQUISITIONS CO., LLC
MUKILTEO, WA 98275



MEDIQUE APAP ACETAMINOPHEN

medique apap acetaminophen tablet, coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51662-1273(NDC:47682-057)
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)		ACETAMINOPHEN	325 mg

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

Product Characteristics			
Color	white	Score	no score
Shape	ROUND	Size	9mm
Flavor		Imprint Code	G323
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51662-1273-2	2 in 1 PACKET; Type 0: Not a Combination Product	03/27/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	03/27/2024	

Labeler - HF Acquisition Co LLC, DBA HealthFirst (045657305)

Registrant - HF Acquisition Co LLC, DBA HealthFirst (045657305)

Establishment			
Name	Address	ID/FEI	Business Operations

HF Acquisition Co LLC, DBA HealthFirst		045657305	repack(51662-1273)
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Revised: 8/2025

HF Acquisition Co LLC, DBA HealthFirst